(FILE 'HOME' ENTERED AT 14:29:59 ON 25 OCT 2001)

=>

	FILE 'BIOSIS, CABA, CAPLUS, EMBASE, LIFESCI, MEDLINE, SCISEARCH,
	USPATFULL, JAPIO' ENTERED AT 14:30:20 ON 25 OCT 2001
L1	15519 S IGG4 OR (IMMUNE GLOBULIN IGG) OR IMMUNOGLOBULIN IGG
L2	1666 S PLASMA AND L1
L3	813 S L2 AND CONCENTRA?
L4	522 S L3 AND PREPAR?
L5	202 S L4 AND LYOPHILIZ?
L6	202 DUP REM L5 (O DUPLICATES REMOVED)
L7	81 S L6 AND IGG4

_

ANSWER 1 OF 81 USPATFULL

2001:184842 USPATFULL ACCESSION NUMBER: Fas antigen derivatives TITLE:

Nakamura, Norio, Tokyo, Japan INVENTOR(S):

Nagata, Shigekazu, Osaka-fu, Japan

PATENT ASSIGNEE(S): Mochida Pharmaceutical Co., Ltd., Tokyo, Japan

(non-U.S. corporation)

Osaka Bioscience Institute, Osaka, Japan (non-U.S.

corporation)

NUMBER KIND DATE US 6306395 B1 20011023 WO 9742319 19971113 PATENT INFORMATION: US 1998-180100 APPLICATION INFO.: 19981102 (9) WO 1997-JP1502 19970501 19981102 PCT 371 date

19981102 PCT 102(e) date

NUMBER · DATE ______

JP 1996-135760 19960502 PRIORITY INFORMATION:

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED PRIMARY EXAMINER: Huff, Sheela ASSISTANT EXAMINER: Harris, Alana M.

LEGAL REPRESENTATIVE: Birch, Stewart, Kolasch & Birch, LLP

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 15 Drawing Figure(s); 28 Drawing Page(s)

LINE COUNT: 2004

ANSWER 2 OF 81 USPATFULL L7

2001:173139 USPATFULL ACCESSION NUMBER: Method of treatment TITLE:

INVENTOR(S): Whitfill, Craig E., Apex, NC, United States Thoma, John A., Fayetteville, AR, United States Fredericksen, Tommy L., Ashford, CT, United States Tyczkowski, Julius K., Cary, NC, United States Thaxton, Jr., J. Paul, Brandon, MS, United States

University of Arkansas, Fayetteville, AR, United States

PATENT ASSIGNEE(S): (U.S. corporation)

NUMBER KIND DATE -----US 6299874 B1 20011009 US 2000-613611 20000711 (9) PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.:

Continuation of Ser. No. US 1998-13760, filed on 27 Jan 1998, now patented, Pat. No. US 6136319 Continuation of Ser. No. US 1996-697268, filed on 21 Aug 1996, now patented, Pat. No. US 5871748 Continuation of Ser. No. US 1994-345291, filed on 28 Nov 1994, now abandoned Continuation of Ser. No. US 1993-8394, filed on 25 Jan 1993, now patented, Pat. No. US 5397569 Continuation of

Ser. No. US 1990-586859, filed on 21 Sep 1990, now abandoned Continuation-in-part of Ser. No. US 1990-480678, filed on 15 Feb 1990, now abandoned Continuation-in-part of Ser. No. US 1989-416035, filed

on 2 Oct 1989, now abandoned

DOCUMENT TYPE: Utility GRANTED FILE SEGMENT:

PRIMARY EXAMINER: Nelson, Brett L.

LEGAL REPRESENTATIVE: Myers Bigel Sibley & Sajovec

NUMBER OF CLAIMS: 116 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

LINE COUNT:

12 Drawing Figure(s); 7 Drawing Page(s)

1982

ANSWER 3 OF 81 USPATFULL

ACCESSION NUMBER:

2001:171199 USPATFULL

TITLE:

Anti-TNF antibodies and peptides of human tumor

necrosis factor

INVENTOR(S):

Le, Junming, Jackson Heights, NY, United States

Vilcek, Jan, New York, NY, United States Daddona, Peter, Menlo Park, CA, United States Ghrayeb, John, Downingtown, PA, United States

Knight, David, Berwyn, PA, United States Siegel, Scott, Westborough, MA, United States

PATENT ASSIGNEE(S):

Centocor, Inc.,, Malvern, PA, United States (U.S.

corporation)

NUMBER KIND DATE ______

PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.: US 2001027249 A1 20011004 US 2001-756301 A1 20010108 20010108 (9)

Division of Ser. No. US 1998-133119, filed on 12 Aug 1998, PENDING Division of Ser. No. US 1995-570674,

filed on 11 Dec 1995, ABANDONED Continuation-in-part of Ser. No. US 1994-324799, filed on 18 Oct 1994, GRANTED, Pat. No. US 5698195 Continuation-in-part of Ser. No. US 1994-192102, filed on 4 Feb 1994, GRANTED, Pat. No. US

5656272 Continuation-in-part of Ser. No. US

1994-192861, filed on 4 Feb 1994, GRANTED, Pat. No. US

5919452 Continuation-in-part of Ser. No. US 1994-192093, filed on 4 Feb 1994, PENDING

Continuation-in-part of Ser. No. US 1993-10406, filed on 29 Jan 1993, ABANDONED Continuation-in-part of Ser. No. US 1993-13413, filed on 2 Feb 1993, ABANDONED

Continuation-in-part of Ser. No. US 1992-943852, filed on 11 Sep 1992, ABANDONED Continuation-in-part of Ser. No. US 1992-853606, filed on 18 Mar 1992, ABANDONED Continuation-in-part of Ser. No. US 1991-670827, filed

on 18 Mar 1991, ABANDONED

DOCUMENT TYPE: FILE SEGMENT:

Utility APPLICATION

LEGAL REPRESENTATIVE:

Carolyn S. Elmore, HAMILTON, BROOK, SMITH & REYNOLDS, P.C., Two Militia Drive, Lexington, MA, 02421-4799

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

1

NUMBER OF DRAWINGS:

37 Drawing Page(s)

LINE COUNT:

5577

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 4 OF 81 USPATFULL

ACCESSION NUMBER:

2001:165585 USPATFULL

TITLE:

INVENTOR(S):

Immunoassay technique using multispecific molecules

Khaw, Ban-an, Milton, MA, United States Narula, Jagat, Rosemont, PA, United States

NUMBER KIND DATE US 2001024795 A1 20010927 US 2000-727421 A1 20001201 (9)

PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.:

Continuation-in-part of Ser. No. US 1999-380168, filed

on 6 Oct 1999, PENDING

DATE NUMBER

PRIORITY INFORMATION: US 1997-39111 19970226 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: PENNIE AND EDMONDS, 1155 AVENUE OF THE AMERICAS, NEW

YORK, NY, 100362711

NUMBER OF CLAIMS: 55 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 3 Drawing Page(s)

LINE COUNT: · 1620

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 5 OF 81 USPATFULL

ACCESSION NUMBER: 2001:165438 USPATFULL

TITLE: Monkey monoclonal antibodies specific to human B7.1

and/or B7.2, primatized forms thereof, pharmaceutical

compositions containing, and use thereof as

immunosuppressants

INVENTOR(S): Anderson, Darrell R., Escondido, CA, United States

Brams, Peter, San Diego, CA, United States

Hanna, Nabil, Rancho Santa Fe, CA, United States Shestowsky, William S., San Diego, CA, United States

Heard, Cheryl, Encinitas, CA, United States

PATENT ASSIGNEE(S): IDEC Pharmaceuticals Corporation (U.S. corporation)

PATENT INFORMATION: US 2001024648 Al 20010927 APPLICATION INFO.: US 2001-758173 Al 20010112 (9)

RELATED APPLN. INFO.: Division of Ser. No. US 1999-383916, filed on 26 Aug 1999, PENDING Division of Ser. No. US 1995-487550, filed on 7 Jun 1995, GRANTED, Pat. No. US 6113898

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: Pillsbury Winthrop LLP, Intellectual Property Group,

East Tower, Ninth Floor, 1100 New York Avenue, N.W.,

Washington, DC, 20005-3918

NUMBER OF CLAIMS: 20 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 22 Drawing Page(s)

LINE COUNT: 1691

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 6 OF 81 USPATFULL

ACCESSION NUMBER: 2001:160973 USPATFULL

TITLE: Use of heregulin as a growth factor

INVENTOR(S): Sliwkowski, Mark X., San Carlos, CA, United States Kern, Jeffrey A., Iowa City, IA, United States

NUMBER KIND DATE

PATENT INFORMATION: US 2001023241 A1 20010920 APPLICATION INFO.: US 2001-773517 A1 20010202 (9)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1999-243198, filed on 2 Feb

1999, ABANDONED

NUMBER DATE

PRIORITY INFORMATION: US 1998-73866 19980204 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: Supervisor, Patent Prosecution Services, PIPER MARBURY

RUDNICK & WOLFE LLP, 1200 Nineteenth Street, N.W.,

Washington, DC, 20036-2412

NUMBER OF CLAIMS:

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 28 Drawing Page(s)

LINE COUNT:

3786

L7 ANSWER 7 OF 81 USPATFULL

ACCESSION NUMBER:

2001:160802 USPATFULL Interleukins-21 and 22

TITLE:

INVENTOR(S):

Ebner, Reinhard, Gaithersburg, MD, United States

Ruben, Steven M., Olney, MD, United States

NUMBER KIND DATE US 2001023070 A1 20010920 US 2000-731816 A1 20001208 (9) PATENT INFORMATION: APPLICATION INFO.:

RELATED APPLN. INFO.:

Continuation-in-part of Ser. No. US 1999-320713, filed on 27 May 1999, PENDING Continuation-in-part of Ser. No. WO 1999-US11644, filed on 27 May 1999, UNKNOWN

NUMBER DATE _____ ___ US 1998-87340 19980529 (60) US 1999-131965 19990430 (60) US 1999-169837 19991209 (60) PRIORITY INFORMATION:

DOCUMENT TYPE: FILE SEGMENT:

Utility APPLICATION

LEGAL REPRESENTATIVE:

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

13 Drawing Page(s)

LINE COUNT:

7740

49

ANSWER 8 OF 81 USPATFULL L7

ACCESSION NUMBER:

2001:155766 USPATFULL 49 human secreted proteins TITLE:

INVENTOR(S):

Moore, Paul A., Germantown, MD, United States Ruben, Steven M., Oley, MD, United States Olsen, Henrik S., Gaithersburg, MD, United States Shi, Yanggu, Gaithersburg, MD, United States Rosen, Craig A., Laytonsville, MD, United States Florence, Kimberly A., Rockville, MD, United States Soppet, Daniel R., Centreville, VA, United States Lafleur, David W., Washington, DC, United States Endress, Gregory A., Potomac, MD, United States Ebner, Reinhard, Gaithersburg, MD, United States Komatsoulis, George, Silver Spring, MD, United States

Duan, Roxanne D., Bethesda, MD, United States

NUMBER KIND DATE US 2001021700 A1 20010913 US 2000-739254 A1 20001219 (9) PATENT INFORMATION:

APPLICATION INFO.: RELATED APPLN. INFO.:

Continuation of Ser. No. US 2000-511554, filed on 23 Feb 2000, ABANDONED Continuation-in-part of Ser. No. WO

1999-US19330, filed on 24 Aug 1999, UNKNOWN

NUMBER DATE

PRIORITY INFORMATION:

US 1998-97917 19980825 (60) US 1998-98634 19980831 (60)

DOCUMENT TYPE: FILE SEGMENT:

Utility APPLICATION

LEGAL REPRESENTATIVE:

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: 23 EXEMPLARY CLAIM: 15462 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 9 OF 81 USPATFULL

2001:147682 USPATFULL ACCESSION NUMBER:

Anti-TNFa antibodies and assays employing anti-TNFa TITLE:

antibodies

Le, Junming, Jackson Heights, NY, United States INVENTOR(S):

> Vilcek, Jan, New York, NY, United States Dadonna, Peter, Palo Alto, CA, United States Ghrayeb, John, Thorndale, PA, United States Knight, David, Berwyn, PA, United States

Siegel, Scott A., Westborough, MA, United States New York University Medical Center, New York, NY,

PATENT ASSIGNEE(S): United States (U.S. corporation)

Centocor, Inc., Malvern, PA, United States (U.S.

corporation)

NUMBER KIND DATE US 6284471 B1 20010904 US 1994-192093 19940204 PATENT INFORMATION:

19940204 (8) APPLICATION INFO.: Continuation-in-part of Ser. No. US 1993-10406, filed RELATED APPLN. INFO.:

on 29 Jan 1993, now abandoned Continuation-in-part of

Ser. No. US 1993-13413, filed on 2 Feb 1993, now abandoned Continuation-in-part of Ser. No. US

1992-943852, filed on 11 Sep 1992, now abandoned Continuation-in-part of Ser. No. US 1992-853606, filed on 18 Mar 1992, now abandoned Continuation-in-part of Ser. No. US 1991-670827, filed on 18 Mar 1991, now

abandoned Utility GRANTED

FILE SEGMENT: PRIMARY EXAMINER: Caputa, Anthony C. ASSISTANT EXAMINER: Canella, Karen A.

Hamilton, Brook, Smith & Reynolds, P.C. LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

DOCUMENT TYPE:

NUMBER OF DRAWINGS: 48 Drawing Figure(s); 36 Drawing Page(s)

LINE COUNT: 5032

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 10 OF 81 USPATFULL L7

2001:142469 USPATFULL ACCESSION NUMBER:

TITLE: Process for producing immunoglobulins for intravenous

administration and other immunoglobulin products

Laursen, Inga, Hellerup, Denmark INVENTOR(S):

Teisner, B.o slashed.rge, Odense C, Denmark Statens Serum Institut, Copenhagen S., Denmark

PATENT ASSIGNEE(S): (non-U.S. corporation)

NUMBER KIND DATE US 6281336 B1 20010828 US 1999-328497 19990609 (9) PATENT INFORMATION: APPLICATION INFO.:

NUMBER DATE ______

EP 1998-201909 19980609 US 1998-102055 19980928 PRIORITY INFORMATION:

19980928 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER:

Saunders, David

LEGAL REPRESENTATIVE:

Birch, Stewart, Kolasch & Birch, LLP

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

14

LINE COUNT:

1 1465

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 11 OF 81 USPATFULL

ACCESSION NUMBER:

2001:139293 USPATFULL

TITLE:

Fibroblast growth factor receptor-5

INVENTOR(S):

Young, Paul E., Gaithersburg, MD, United States

Ruben, Steven M., Olney, MD, United States

NUMBER KIND DATE ______

PATENT INFORMATION: APPLICATION INFO.:

US 2001016335 A1 20010823 US 2001-758386 A1 20010112 (9)

RELATED APPLN. INFO.:

Continuation of Ser. No. US 1999-293182, filed on 16

Apr 1999, ABANDONED

NUMBER DATE -----

PRIORITY INFORMATION:

US 1998-105465 19981023 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS:

23

EXEMPLARY CLAIM:

1

NUMBER OF DRAWINGS:

10 Drawing Page(s)

LINE COUNT:

6097

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 12 OF 81 USPATFULL

ACCESSION NUMBER:

2001:136770 USPATFULL

TITLE:

Anti-TNF antibodies and peptides of human tumor

necrosis factor

INVENTOR(S):

Le, Junming, Jackson Heights, NY, United States Vilcek, Jan, New York, NY, United States Daddona, Peter, Menlo Park, CA, United States Ghrayeb, John, Thorndale, PA, United States Knight, David, Berwyn, PA, United States Siegel, Scott, Westborough, MA, United States

PATENT ASSIGNEE(S): New York University, New York, NY, United States (U.S.

corporation)

Centocor, Inc., Malvern, PA, United States (U.S.

corporation)

New York University Medical Center, New York, NY,

United States (U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: APPLICATION INFO.:

US 6277969 B1 US 1998-133119 В1 20010821 19980812 (9)

RELATED APPLN. INFO.:

Division of Ser. No. US 1995-570674, filed on 11 Dec 1995, now abandoned Continuation-in-part of Ser. No. US 1994-324799, filed on 18 Oct 1994, now patented, Pat.

No. US 5698195, issued on 16 Dec 1997

Continuation-in-part of Ser. No. US 1994-192102, filed on 4 Feb 1994, now patented, Pat. No. US 5656272,

issued on 12 Aug 1997 Continuation-in-part of Ser. No. US 1994-192861, filed on 4 Feb 1994, now patented, Pat. No. US 5919452, issued on 6 Jul 1999

Continuation-in-part of Ser. No. US 1994-192093, filed

on 4 Feb 1994 Continuation-in-part of Ser. No. US 1993-10406, filed on 29 Jan 1993, now abandoned Continuation-in-part of Ser. No. US 1993-13413, filed on 2 Feb 1993, now abandoned Continuation-in-part of Ser. No. US 1992-943852, filed on 11 Sep 1992, now abandoned Continuation-in-part of Ser. No. US 1992-853606, filed on 18 Mar 1992, now abandoned Continuation-in-part of Ser. No. US 1991-670827, filed

on 18 Mar 1991, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Caputa, Anthony C. ASSISTANT EXAMINER: Canella, Karen

LEGAL REPRESENTATIVE: Hamilton, Brook, Smith & Reynolds, P.C.

NUMBER OF CLAIMS: 4 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 49 Drawing Figure(s); 37 Drawing Page(s)

LINE COUNT: 5429

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 13 OF 81 USPATFULL

ACCESSION NUMBER: 2001:136184 USPATFULL

TITLE: Immunoglobulin-like domains with increased half-lives INVENTOR(S): Ward, Elizabeth Sally, Dallas, TX, United States PATENT ASSIGNEE(S): Board of Regents, The University of Texas System,

Austin, TX, United States (U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 6277375 B1 20010821 APPLICATION INFO.: US 1997-811463 19970303 (8)

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Saunders, David LEGAL REPRESENTATIVE: Fulbright & Jaworski

NUMBER OF CLAIMS: 4
EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 19 Drawing Figure(s); 15 Drawing Page(s)

LINE COUNT: 4495

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 14 OF 81 USPATFULL

ACCESSION NUMBER: 2001:102968 USPATFULL

TITLE: High affinity human antibodies and human antibodies

against digoxin

INVENTOR(S): Lonberg, Nils, Woodside, CA, United States

Kay, Robert M., San Francisco, CA, United States

PATENT ASSIGNEE(S): GenPharm International, San Jose, CA, United States

(U.S. corporation)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1996-758417, filed

on 2 Dec 1996 Continuation-in-part of Ser. No. US 1996-728463, filed on 10 Oct 1996 Continuation-in-part of Ser. No. US 1995-544404, filed on 10 Oct 1995, now patented, Pat. No. US 5770429 Continuation-in-part of Ser. No. US 1994-352322, filed on 7 Dec 1994, now patented, Pat. No. US 5625126 Continuation-in-part of Ser. No. US 1994-209741, filed on 9 Mar 1994, now

abandoned Continuation-in-part of Ser. No. US 1993-165699, filed on 10 Dec 1993, now abandoned

Continuation-in-part of Ser. No. US 1993-161739, filed on 3 Dec. 1993, now abandoned Continuation-in-part of Ser. No. US 1993-155301, filed on 18 Nov 1993, now abandoned Continuation-in-part of Ser. No. US 1993-96762, filed on 22 Jul 1993, now patented, Pat. No. US 5814318 Continuation-in-part of Ser. No. US 1993-53131, filed on 26 Apr 1993, now patented, Pat. No. US 5661016 Continuation-in-part of Ser. No. US 1992-990860, filed on 16 Dec 1992, now patented, Pat. No. US 5545806 Continuation-in-part of Ser. No. US 1992-904068, filed on 23 Jun 1992, now abandoned Continuation-in-part of Ser. No. US 1992-853408, filed on 18 Mar 1992, now patented, Pat. No. US 5789650 Continuation-in-part of Ser. No. US 1992-834539, filed on 5 Feb 1992, now patented, Pat. No. US 5633425 Continuation-in-part of Ser. No. US 1991-810279, filed on 17 Dec 1991, now patented, Pat. No. US 5569825 Continuation-in-part of Ser. No. US 1990-575962, filed on 31 Aug 1990, now abandoned Continuation-in-part of Ser. No. US 1990-574748, filed on 29 Aug 1990, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Chan, Christina Y. ASSISTANT EXAMINER: DiBrino, Marianne

LEGAL REPRESENTATIVE: Townsend and Townsend and Crew LLP

NUMBER OF CLAIMS: 2 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 119 Drawing Figure(s); 103 Drawing Page(s)

LINE COUNT: 10059

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 15 OF 81 USPATFULL

ACCESSION NUMBER: 2001:97420 USPATFULL

TITLE: Methods of inhibiting inflammation at the site of a

central nervous system injury with alphaD-specific

antibodies

INVENTOR(S): Gallatin, W. Michael, 8412 SE. 33rd Pl., Mercer Island,

WA, United States 98040

Van der Vieren, Monica, 2446 NW. 64th St., Seattle, WA,

United States 98107

PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.:

Continuation-in-part of Ser. No. US 1997-943363, filed on 3 Oct 1997, now patented, Pat. No. US 5837478, issued on 17 Nov 1998 Continuation-in-part of Ser. No. US 1996-605672, filed on 22 Feb 1996, now patented, Pat. No. US 5817515, issued on 6 Oct 1998

Continuation-in-part of Ser. No. US 1994-362652, filed on 21 Dec 1994, now patented, Pat. No. US 5766850, issued on 16 Jun 1998 Continuation-in-part of Ser. No.

US 1994-286889, filed on 5 Aug 1994, now patented, Pat. No. US 5470953, issued on 28 Nov 1995

Continuation-in-part of Ser. No. US 1993-173497, filed on 23 Dec 1993, now patented, Pat. No. US 5437958,

issued on 1 Aug 1995

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Gambel, Phillip

NUMBER OF CLAIMS: 10 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT: 6697

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 16 OF 81 USPATFULL

ACCESSION NUMBER: 2001:67171 USPATFULL

TITLE: Antithrombotic agent and humanized anti-von Willebrand

factor monoclonal antibody

INVENTOR(S): Co, Man Sung, Cupertino, CA, United States

Vasquez, Maximiliano, Palo Alto, CA, United States

PATENT ASSIGNEE(S): Ajinomoto Co., Inc., Tokyo, Japan (non-U.S.

corporation)

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Gambel, Phillip

LEGAL REPRESENTATIVE: Oblon, Spivak, McClelland, Maier & Neustadt, P.C.

NUMBER OF CLAIMS: 9 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 784

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 17 OF 81 USPATFULL

ACCESSION NUMBER: 2001:47546 USPATFULL

TITLE: Humanized antibodies reactive with L-selectin INVENTOR(S): Co, Man Sung, Cupertino, CA, United States

PATENT ASSIGNEE(S): Protein Design Labs, Inc., Fremont, CA, United States

(U.S. corporation)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1993-160074, filed

on 30 Nov 1993, now abandoned Continuation-in-part of Ser. No. US 1992-983946, filed on 1 Dec 1992, now

abandoned

DOCUMENT TYPE: Utility
FILE SEGMENT: Granted

PRIMARY EXAMINER: Gambel, Phillip

LEGAL REPRESENTATIVE: Townsend & Townsend & Crew

NUMBER OF CLAIMS: 17 EXEMPLARY CLAIM: 1,13,15

NUMBER OF DRAWINGS: 16 Drawing Figure(s); 13 Drawing Page(s)

LINE COUNT: 1947

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 18 OF 81 USPATFULL

ACCESSION NUMBER: 2001:47545 USPATFULL

TITLE: Cross-reacting monoclonal antibodies specific for

E-selectin and P-selectin

INVENTOR(S): Berg, Ellen L., Palo Alto, CA, United States

PATENT ASSIGNEE(S): Protein Design Labs, Inc., Fremont, CA, United States

(U.S. corporation)

NUMBER KIND DATE US 6210670 B1 20010403 WO 9534324 19951221 PATENT INFORMATION: APPLICATION INFO.: US 1996-619491 19960326 WO 1995-US7302 19950607 19960326 PCT 371 date

19960326 PCT 102(e) date

Continuation-in-part of Ser. No. US 1994-259963, filed RELATED APPLN. INFO.:

on 14 Jun 1994, now patented, Pat. No. US 5622701

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

Gambel, Phillip PRIMARY EXAMINER:

Townsend & Townsend & Crew LLP, Storella, John LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: 5.5 EXEMPLARY CLAIM: 1,15

NUMBER OF DRAWINGS: 20 Drawing Figure(s); 14 Drawing Page(s)

LINE COUNT: 1987

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 19 OF 81 USPATFULL

2001:4887 USPATFULL ACCESSION NUMBER:

Anti-IgE antibodies and method of improving TITLE:

polypeptides

Lowman, Henry B., El Granada, CA, United States INVENTOR(S):

Presta, Leonard G., San Francisco, CA, United States Jardieu, Paula M., San Mateo, CA, United States

Lowe, John, Daly City, CA, United States

Genentech, Inc., South San Francisco, CA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE _____ ___ US 6172213 US 1998-109207 В1 PATENT INFORMATION: 20010109 APPLICATION INFO.: 19980630 (9)

> NUMBER DATE _____

US 1997-51554 19970702 (60) PRIORITY INFORMATION:

DOCUMENT TYPE: Patent FILE SEGMENT: Granted

PRIMARY EXAMINER: Chan, Christina Y. ASSISTANT EXAMINER: Ewoldt, Gerald R. LEGAL REPRESENTATIVE: Svoboda, Craig G.

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 23 Drawing Figure(s); 19 Drawing Page(s)

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 20 OF 81 USPATFULL

2000:168146 USPATFULL ACCESSION NUMBER:

Anti-human .alpha..sub.v .beta..sub.3 and .alpha..sub.v TITLE:

.beta..sub.5 antibodies

INVENTOR(S): Jonak, Zdenka Ludmila, SmithKline Beecham Corporation Corporate Intellectual Property-UW2220 P.O. Box 1539,

King of Prussia, PA, United States 19406-0939 Taylor, Alexander, SmithKline Beecham Corporation Corporate Intellectual Property-UW2220 P.O. Box 1539,

King of Prussia, PA, United States 19406-0939 Trulli, Stephen H, SmithKline Beecham Corporation Corporate Intellectual Property-UW2220 P.O. Box 1539,

King of Prussia, PA, United States 19406-0939

Johanson, Kyung O, SmithKline Beecham Corporation Corporate Intellectual Property-UW2220 P.O. Box 1539, King of Prussia, PA, United States 19406-0939

NUMBER KIND DATE

PATENT INFORMATION: APPLICATION INFO.: US 6160099 20001212 US 1998-199149 19981124 (9)

DOCUMENT TYPE: Utility Granted FILE SEGMENT: PRIMARY EXAMINER: PRIMARY EXAMINER: Huff, Sheela ASSISTANT EXAMINER: Helms, Larry R.

LEGAL REPRESENTATIVE: Baumeister, Kirk, King, William T.

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

12 Drawing Figure(s); 9 Drawing Page(s) 2245 NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 21 OF 81 USPATFULL

ACCESSION NUMBER: 2000:167510 USPATFULL Uses of Wnt polypeptides TITLE:

Matthews, William, Woodside, CA, United States INVENTOR(S): Austin, Timothy W., Morgan Hill, CA, United States

Genentech, Inc., So. San Francisco, CA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE _____ __ US 6159462 20001212 US 1997-911860 19970815 PATENT INFORMATION: APPLICATION INFO.: 19970815 (8)

NUMBER DATE _____

US 1996-24068 19960816 (60) PRIORITY INFORMATION:

DOCUMENT TYPE: Utility FILE SEGMENT: Granted PRIMARY EXAMINER:

PRIMARY EXAMINER: Saunders, David
ASSISTANT EXAMINER: VanderVegt, F. Pierre

LEGAL REPRESENTATIVE: Svoboda, Craig G., Carpenter, David A.

NUMBER OF CLAIMS: 13 EXEMPLARY CLAIM:

INVENTOR(S):

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 2 Drawing Page(s)

3907 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 22 OF 81 USPATFULL

ACCESSION NUMBER: 2000:157557 USPATFULL

TITLE: Monoclonal antibodies specific for the extracellular

> domain of prostate-specific membrane antigen Murphy, Gerald P., Seattle, WA, United States Boynton, Alton L., Redmond, WA, United States

Holmes, Eric H., Bothell, WA, United States Tino, William Thomas, Redmond, WA, United States

Northwest Biotherapeutics, Inc., Seattle, WA, United PATENT ASSIGNEE(S):

States (U.S. corporation)

NUMBER KIND DATE -----US 6150508 US 1998-44668 PATENT INFORMATION: 20001121 APPLICATION INFO.: 19980318 (9)

Continuation-in-part of Ser. No. US 1997-827017, filed RELATED APPLN. INFO.:

on 25 Mar 1997, now abandoned which is a

continuation-in-part of Ser. No. US 1996-621399, filed

on 25 Mar 1996, now abandoned

DOCUMENT TYPE:

Utility Granted FILE SEGMENT:

PRIMARY EXAMINER:

Ungar, Susan

LEGAL REPRESENTATIVE:

Townsend and Townsend and Crew LLP

NUMBER OF CLAIMS:

16

EXEMPLARY CLAIM:

1,2,7,12

NUMBER OF DRAWINGS:

24 Drawing Figure(s); 20 Drawing Page(s)

LINE COUNT:

1896

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 23 OF 81 USPATFULL

ACCESSION NUMBER:

PATENT ASSIGNEE(S):

2000:146146 USPATFULL

TITLE:

Cell-targeting molecule comprising a mutant human

carboxypeptidase A

INVENTOR(S):

Smith, Gary Keith, Raleigh, NC, United States

Blumenkopf, Todd Andrew, Old Lyme, CT, United States

Cory, Michael, Chapel Hill, NC, United States

Glaxo Wellcome Inc., Research Triangle Park, NC, United

States (U.S. corporation)

	NUMBER	KIND DATE	
PATENT INFORMATION:	US 6140100	20001031	
	WO 9513095	19950518	
APPLICATION INFO.:	US 1996-640906	19960509	(8)
	WO 1994-GB2483	19941111	
		19960509	PCT 371 date
		19960509	PCT 102(e) date

NUMBER	DATE

PRIORITY INFORMATION:

GB 1993-23429 19931112 Utility

DOCUMENT TYPE: FILE SEGMENT:

Granted

PRIMARY EXAMINER:

Achutamurthy, Ponnathapura

ASSISTANT EXAMINER:

Moore, William W.

LEGAL REPRESENTATIVE:

Grassler, Frank P., Bennett, Virginia C., Hrubiec,

Robert T.

NUMBER OF CLAIMS:

12

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

7 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT:

7473

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 24 OF 81 USPATFULL 1.7

ACCESSION NUMBER:

2000:141887 USPATFULL

TITLE: INVENTOR(S):

Method of treating viral diseases in animals Whitfill, Craig E., Apex, NC, United States Thoma, John A., Fayetteville, AR, United States Fredericksen, Tommy L., Ashford, CT, United States Tyczkowski, Julius K., Cary, NC, United States Thaxton, Jr., J. Paul, Brandon, MS, United States

PATENT ASSIGNEE(S):

The University of Arkansas, Fayetteville, AR, United

States (U.S. corporation)

	NUMBER	KIND	DATE	
PATENT INFORMATION:	US 6136319		20001024	
APPLICATION INFO.:	US 1998-13760		19980127	(9)

RELATED APPLN. INFO.:

Continuation of Ser. No. US 1996-697268, filed on 21 Aug 1996, now patented, Pat. No. US 5871748 which is a continuation of Ser. No. US 1994-345291, filed on 28 Nov 1994, now abandoned which is a continuation of Ser. No. US 1993-8394, filed on 25 Jan 1993, now patented, Pat. No. US 5397569 which is a continuation of Ser. No. US 1990-586859, filed on 21 Sep 1990, now abandoned

which is a continuation-in-part of Ser. No. US

1990-480678, filed on 15 Feb 1990, now abandoned which is a continuation-in-part of Ser. No. US 1989-416035,

filed on 2 Oct 1989, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Stucker, Jeffrey Nelson, Brett ASSISTANT EXAMINER:

Myers Bigel Sibley & Sajovec LEGAL REPRESENTATIVE: NUMBER OF CLAIMS: 28

EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 12 Drawing Figure(s); 7 Drawing Page(s)

1766 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 25 OF 81 USPATFULL

2000:141878 USPATFULL ACCESSION NUMBER:

Recombinant anti-CD4 antibodies for human therapy TITLE:

INVENTOR(S): Hanna, Nabil, Olivenhain, CA, United States

Newman, Roland Anthony, San Diego, CA, United States Reff, Mitchell Elliot, San Diego, CA, United States IDEC Pharmaceuticals Corporation, San Diego, CA, United

PATENT ASSIGNEE(S):

States (U.S. corporation)

NUMBER _____ ___

US 6136310 US 1995-523894 20001024 PATENT INFORMATION: APPLICATION INFO.: 19950906

Continuation-in-part of Ser. No. US 1995-476237, filed RELATED APPLN. INFO.: on 7 Jun 1995, now patented, Pat. No. US 5756096 which is a continuation-in-part of Ser. No. US 1995-379072,

filed on 25 Jan 1995, now patented, Pat. No. US 5658570 which is a continuation of Ser. No. US 1992-912292, filed on 10 Jul 1992, now abandoned which is a

continuation-in-part of Ser. No. US 1992-856281, filed

on 23 Mar 1992, now abandoned which is a

continuation-in-part of Ser. No. US 1991-735064, filed

on 25 Jul 1991, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

Bansal, Geetha P. PRIMARY EXAMINER:

Burns, Doane, Swecker & Mathis, LLP LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: 16 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 32 Drawing Figure(s); 32 Drawing Page(s)

LINE COUNT: 3398

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 26 OF 81 USPATFULL L7

ACCESSION NUMBER: 2000:124800 USPATFULL

Altered polypeptides with increased half-life TITLE:

Presta, Leonard G., San Francisco, CA, United States INVENTOR(S): Snedecor, Bradley R., Portola Valley, CA, United States

Genentech, Inc., S. San Francisco, CA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE ______ US 6121022 US 1995-422112 20000919 PATENT INFORMATION: 19950414 (8) APPLICATION INFO.:

DOCUMENT TYPE: Utility FILE SEGMENT:

Granted

PRIMARY EXAMINER: Saunders, David

LEGAL REPRESENTATIVE: Lee, WendyFlehr Hohabch Test Albritton & Herbert LLP

NUMBER OF CLAIMS:

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

4 Drawing Figure(s); 3 Drawing Page(s)

LINE COUNT:

3411

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 27 OF 81 USPATFULL

ACCESSION NUMBER:

2000:117279 USPATFULL

TITLE:

Human B7.1-specific primatized antibodies and

transfectomas expressing said antibodies

INVENTOR(S):

Anderson, Darrell R., Escondido, CA, United States

Brams, Peter, San Diego, CA, United States

Hanna, Nabil, Rancho Santa Fe, CA, United States Shestowsky, William S., San Diego, CA, United States

Heard, Cheryl, Encinitas, CA, United States

PATENT ASSIGNEE(S):

IDEC Pharmaceuticals Corporation, San Diego, CA, United

States (U.S. corporation)

NUMBER KIND DATE _____

PATENT INFORMATION:
APPLICATION INFO.:

US 6113898 20000905 US 1995-487550 19950607 (8)

DOCUMENT TYPE: Utility FILE SEGMENT: Granted PRIMARY EXAMINER: PRIMARY EXAMINER: Feisee, Lila
ASSISTANT EXAMINER: Gambel, Phillip

LEGAL REPRESENTATIVE:

Burns, Doane, Swecker & Mathis, LLP

NUMBER OF CLAIMS:

11

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

22 Drawing Figure(s); 22 Drawing Page(s)

LINE COUNT:

2309

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

1.7 ANSWER 28 OF 81 USPATFULL

ACCESSION NUMBER:

2000:98553 USPATFULL

TITLE:

Polypeptides altered to contain an epitope from the Fc

region of an IgG molecule for increased half-life Presta, Leonard G., San Francsico, CA, United States INVENTOR(S):

Snedecor, Bradley R., Portola Valley, CA, United States Genentech, Inc., S. San Francisco, CA, United States

PATENT ASSIGNEE(S): (U.S. corporation)

> NUMBER KIND DATE -----

PATENT INFORMATION: APPLICATION INFO.:

US 6096871 20000801 US 1995-422093 19950414 (8)

DOCUMENT TYPE:

FILE SEGMENT:

Utility Granted

FILE SEGMENT: Granted PRIMARY EXAMINER: Reeves, Julie

LEGAL REPRESENTATIVE: Hasak, Jan, Vance, Dolly A. Flehr Hohbach Test Albritton

& Herbert LLP

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

14 1

NUMBER OF DRAWINGS:

4 Drawing Figure(s); 3 Drawing Page(s)

LINE COUNT:

3391

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 29 OF 81 USPATFULL L.7

ACCESSION NUMBER:

2000:70963 USPATFULL

TITLE:

Reshaped monoclonal antibodies against an

immunoglobulin isotype

Hardman, Norman, Riehen, Switzerland INVENTOR(S):

Kolbinger, Frank, Freiburg, Germany, Federal Republic

of

Saldanha, Jose, Enfield, United Kingdom

Novartis Corporation, Summit, NJ, United States (U.S. PATENT ASSIGNEE(S):

corporation)

Tanox Biosystems, Inc., Houston, TX, United States

(U.S. corporation)

NUMBER KIND DATE _____ -----

PATENT INFORMATION:

APPLICATION INFO .: RELATED APPLN. INFO.: US 6072035 20000606 US 1995-485246 19950607 (8) Division of Ser. No. US 1993-127721, filed on 27 Sep

1993 which is a continuation-in-part of Ser. No. US 1992-952802, filed on 25 Sep 1992, now abandoned

> Utility Granted

> > 20

FILE SEGMENT: Granted PRIMARY EXAMINER: Reeves, Julie LEGAL REPRESENTATIVE:

DOCUMENT TYPE:

FILE SEGMENT:

Ferraro, Gregory D.

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

2 Drawing Figure(s); 1 Drawing Page(s)

LINE COUNT: 2732

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 30 OF 81 USPATFULL

ACCESSION NUMBER:

2000:64943 USPATFULL

TITLE:

Reshaped monoclonal antibodies against an

immunoglobulin isotype

INVENTOR(S):

Hardman, Norman, Riehen, Switzerland

Kolbinger, Frank, Freiburg, Germany, Federal Republic

of

Saldanha, Jose, Enfield, United Kingdom

PATENT ASSIGNEE(S):

Novartis Corporation, Summit, NJ, United States (U.S.

corporation)

Tanox Biosystems, Inc., Houston, TX, United States

(U.S. corporation)

NUMBER KIND DATE _____ -_----

PATENT INFORMATION: US 6066718 20000523 APPLICATION INFO.: US 1993-127721 19930927 (8)

RELATED APPLN. INFO.:

Continuation-in-part of Ser. No. US 1992-952802, filed

on 25 Sep 1992, now abandoned

DOCUMENT TYPE: FILE SEGMENT:

Utility Granted

PRIMARY EXAMINER: Hutzell, Paula K.
ASSISTANT EXAMINER: Worrall, Timothy A. LEGAL REPRESENTATIVE: Ferrar, Gregory D.

NUMBER OF CLAIMS:

10

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

1 1 Drawing Figure(s); 1 Drawing Page(s)

LINE COUNT:

2830

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 31 OF 81 USPATFULL 1.7

ACCESSION NUMBER:

2000:46884 USPATFULL

TITLE:

Blockade of T lymphocyte down-regulation associated

with CTLA-4 signaling

INVENTOR(S):

Allison, James Patrick, Berkeley, CA, United States

Leach, Dana R., Albany, CA, United States

Krummel, Matthew F., Berkeley, CA, United States

PATENT ASSIGNEE(S): The Regents of the University of California, Office of Technology Transfer, Oakland, CA, United States (U.S. corporation)

NUMBER KIND DATE US 6051227 20000418 US 1996-760288 19961204 (8) PATENT INFORMATION: APPLICATION INFO.:

Continuation-in-part of Ser. No. US 1996-646605, filed RELATED APPLN. INFO.:

on 8 May 1996, now patented, Pat. No. US 5811097 which is a continuation-in-part of Ser. No. US 1995-566853, filed on 4 Dec 1995, now patented, Pat. No. US 5855887

which is a continuation-in-part of Ser. No. US 1995-506666, filed on 25 Jul 1995, now abandoned

DOCUMENT TYPE: Utility Granted FILE SEGMENT:

Scheiner, Toni R. PRIMARY EXAMINER:

Trecartin, Richard F., Lorenz, Todd A.Flehr Hohbach LEGAL REPRESENTATIVE:

Test Albritton & Herbert LLP

NUMBER OF CLAIMS: 23 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 33 Drawing Figure(s); 16 Drawing Page(s)

LINE COUNT: 2146

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 32 OF 81 USPATFULL

2000:27563 USPATFULL ACCESSION NUMBER:

Method for detecting the presence of P-selectin TITLE: Chesnut, Robert W., Cardiff, CA, United States INVENTOR(S): Polley, Margaret J., La Jolla, CA, United States Paulson, James C., Del Mar, CA, United States Jones, S. Tarran, Radlett, United Kingdom Saldanha, Jose W., Middlesex, United Kingdom

Bendig, Mary M., London, United Kingdom

Kriegler, Michael, Rancho Santa Fe, CA, United States

Perez, Carl, San Diego, CA, United States Bayer, Robert, San Diego, CA, United States Nunn, Michael, San Diego, CA, United States

Cytel Corporation, San Diego, CA, United States (U.S. PATENT ASSIGNEE(S):

corporation)

NUMBER KIND DATE PATENT INFORMATION: US 6033667 20000307 US 1997-964690 19971105 (8)

APPLICATION INFO.:

Continuation of Ser. No. US 1994-202047, filed on 25 RELATED APPLN. INFO.: Feb 1994, now patented, Pat. No. US 5800815 which is a continuation-in-part of Ser. No. US 1993-57292, filed

on 5 May 1993, now abandoned which is a

continuation-in-part of Ser. No. US 1992-880196, filed

on 5 May 1992, now abandoned

NUMBER DATE

PRIORITY INFORMATION: IL 1993-105614 19930505

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

Chan, Christina Y. PRIMARY EXAMINER: ASSISTANT EXAMINER: Gambel, Phillip LEGAL REPRESENTATIVE: Campbell & Flores LLP

NUMBER OF CLAIMS: 21 EXEMPLARY CLAIM: 1

44 Drawing Figure(s); 40 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT: 4009

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 33 OF 81 USPATFULL

2000:12616 USPATFULL ACCESSION NUMBER: Chimeric antibodies TITLE:

Hardman, Norman, Riehen, Switzerland INVENTOR(S):

Gill, Laura Lee, Riehen, Switzerland

de Winter, Ronald F. J., Milton Ernest, United Kingdom

Wagner, Kathrin, Basel, Switzerland

Heusser, Christoph, Bottmingen, Switzerland

Ciba-Geigy Corporation, Tarrytown, NY, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE US 6020153 20000201 US 1994-307087 19940916 (8) PATENT INFORMATION: APPLICATION INFO.:

Continuation of Ser. No. US 1992-947897, filed on 18 RELATED APPLN. INFO.: Sep 1992, now abandoned which is a continuation of Ser.

No. US 1988-287178, filed on 21 Dec 1988, now abandoned

NUMBER DATE GB 1988-77 19880105 GB 1988-20099 19880824 PRIORITY INFORMATION:

DOCUMENT TYPE: Utility

FILE SEGMENT:

PRIMARY EXAMINER:

Ziska, Suzanne E.

Nowak, Henry P., Elmer, James Scott, Foley, Shawn P.

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 7 Drawing Figure(s); 7 Drawing Page(s)

LINE COUNT: 2592

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 34 OF 81 USPATFULL L7

ACCESSION NUMBER: 2000:10021 USPATFULL

TITLE: Antibody against human interleukin-5-receptor .alpha.

chain

Koike, Masamichi, Tokyo, Japan INVENTOR(S): Furuya, Akiko, Tokyo, Japan

Nakamura, Kazuyasu, Tokyo, Japan Iida, Akihiro, Tokyo, Japan Anazawa, Hideharu, Tokyo, Japan Hanai, Nobuo, Kanagawa, Japan Takatsu, Kiyoshi, Tokyo, Japan

Kyowa Hakko Kogyo Co., Ltd., Tokyo, Japan (non-U.S. PATENT ASSIGNEE(S):

corporation)

NUMBER KIND DATE ______ US 6018032 WO 9710354 PATENT INFORMATION: 20000125 19970320 US 1997-836561 19970509 (8) APPLICATION INFO.: WO 1996-JP2588 19960911 19970509 PCT 371 date

19970509 PCT 102(e) date

NUMBER DATE ______ JP 1995-232384 19950911 PRIORITY INFORMATION:

DOCUMENT TYPE: Utility Granted FILE SEGMENT: PRIMARY EXAMINER: Granted Mertz, Prema

LEGAL REPRESENTATIVE: Pennie & Edmonds LLP

13 NUMBER OF CLAIMS: EXEMPLARY CLAIM:

61 Drawing Figure(s); 61 Drawing Page(s)
5703 NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 35 OF 81 USPATFULL

1999:163215 USPATFULL ACCESSION NUMBER:

Humanized antibodies to human gp39, compositions TITLE:

containing thereof

Black, Amelia, Cardiff, CA, United States INVENTOR(S):

Hanna, Nabil, Olivenhian, CA, United States Padlan, Eduardo A., Kensington, MD, United States Newman, Roland A., San Diego, CA, United States

Idec Pharmaceuticals Corporation, San Diego, CA, United PATENT ASSIGNEE(S):

States (U.S. corporation)

NUMBER KIND DATE _______

US 6001358 19991214 US 1995-554840 19951107 (8) PATENT INFORMATION:
APPLICATION INFO.:

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Chan, Christina Y. ASSISTANT EXAMINER: Gambel, Phillip

LEGAL REPRESENTATIVE: Burns, Doane, Swecker & Mathis, L.L.P.

NUMBER OF CLAIMS: 12 EXEMPLARY CLAIM: 1

15 Drawing Figure(s); 15 Drawing Page(s) NUMBER OF DRAWINGS:

2693 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 36 OF 81 USPATFULL

ACCESSION NUMBER: 1999:155894 USPATFULL

Anti-IgE antibodies and methods of improving TITLE:

polypeptides

Lowman, Henry B., El Granada, CA, United States INVENTOR(S):

Presta, Leonard G., San Francisco, CA, United States

Jardieu, Paula M., San Mateo, CA, United States

Lowe, John, Daly City, CA, United States

Genentech, Inc., South San Francisco, CA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE _____ PATENT INFORMATION: US 5994511 19991130
APPLICATION INFO.: US 1997-887352 19970702 (8)

DOCUMENT TYPE: Utility

FILE SEGMENT: Granted
PRIMARY EXAMINER: Saunders, David LEGAL REPRESENTATIVE: Svoboda, Craig G.

NUMBER OF CLAIMS: 11 1 EXEMPLARY CLAIM:

21 Drawing Figure(s); 19 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT: 5816

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 37 OF 81 USPATFULL L7

1999:136684 USPATFULL ACCESSION NUMBER:

Inhibition of intimal hyperplasia using antibodies to TITLE:

PDGF receptors and heparin

Hart, Charles E., Brier, WA, United States INVENTOR(S):

Kenagy, Richard D., Seattle, WA, United States Clowes, Alexander W., Seattle, WA, United States

ZymoGenetics, Inc., Seattle, WA, United States (U.S. PATENT ASSIGNEE(S):

corporation)

US 5976534 19991102 US 1995-482533 19950607 (8) PATENT INFORMATION: APPLICATION INFO.:

Continuation-in-part of Ser. No. US 1994-366860, filed RELATED APPLN. INFO.:

NUMBER KIND DATE

on 30 Dec 1994, now patented, Pat. No. US 5620687 which is a continuation-in-part of Ser. No. US 1994-304623,

filed on 12 Sep 1994, now abandoned which is a continuation of Ser. No. US 1993-23504, filed on 25 Feb

1993, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted Feisee, Lila PRIMARY EXAMINER: ASSISTANT EXAMINER: Gambel, Phillip Parker, Gary E. LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: 37 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 12 Drawing Figure(s); 12 Drawing Page(s)

LINE COUNT: 2864

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 38 OF 81 USPATFULL

1999:117280 USPATFULL ACCESSION NUMBER:

TITLE: Reshaped monoclonal antibodies against an

immunoglobulin isotype

INVENTOR(S): Hardman, Norman, Riehen, Switzerland

Kolbinger, Frank, Freiburg, Germany, Federal Republic

οf

Saldanha, Jose, Enfield, United Kingdom

Novartis Corporation, Summit, NJ, United States (U.S. PATENT ASSIGNEE(S):

corporation)

Tanox Biosystems, Inc., Houston, TX, United States

(U.S. corporation)

NUMBER KIND DATE -----PATENT INFORMATION: US 5958708 19990928 US 1995-476176 19950607 (8)

APPLICATION INFO.:

Division of Ser. No. US 1993-127721, filed on 27 Sep RELATED APPLN. INFO.: 1993 which is a continuation-in-part of Ser. No. US 1992-952802, filed on 25 Sep 1992, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Caputa, Anthony C. ASSISTANT EXAMINER: Navarro, Mark LEGAL REPRESENTATIVE: Ferraro, Gregory D.

NUMBER OF CLAIMS: 8 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 1 Drawing Figure(s); 1 Drawing Page(s)

LINE COUNT: 2666

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 39 OF 81 USPATFULL

ACCESSION NUMBER: 1999:85255 USPATFULL

TITLE: DNA encoding recombinant IL4 antibodies useful in

treatment of IL4 mediated disorders

INVENTOR(S): Holmes, Stephen D., Epsom, United Kingdom

Gross, Mitchell Stuart, Wayne, PA, United States Sylvester, Daniel R., Phoenixville, PA, United States

PATENT ASSIGNEE(S): SmithKline Beecham Corporation, Philadelphia, PA,

United States (U.S. corporation)

SmithKline Beecham P.L.C., Brentford, United Kingdom (non-U.S. corporation)

NUMBER KIND DATE US 5928904 19990727 US 1995-483632 19950607 (8) PATENT INFORMATION: APPLICATION INFO.: Continuation-in-part of Ser. No. WO 1994-US10308, filed RELATED APPLN. INFO .: on 7 Sep 1994 which is a continuation-in-part of Ser. No. US 1993-136783, filed on 14 Oct 1993, now abandoned which is a continuation of Ser. No. US 1993-117366, filed on 7 Sep 1993, now abandoned DOCUMENT TYPE: Utility FILE SEGMENT: Granted Caputa, Anthony C. PRIMARY EXAMINER: Navarro, Mark ASSISTANT EXAMINER: Eagle, Alissa M., King, William T., Venetianer, Stephen LEGAL REPRESENTATIVE: NUMBER OF CLAIMS: 33 EXEMPLARY CLAIM: 1 12 Drawing Figure(s); 11 Drawing Page(s) NUMBER OF DRAWINGS: LINE COUNT: 2462 CAS INDEXING IS AVAILABLE FOR THIS PATENT. ANSWER 40 OF 81 USPATFULL 1999:81543 USPATFULL ACCESSION NUMBER: Soluble lymphotoxin-.beta. receptors and TITLE: anti-lymphotoxin receptor and ligand antibodies as therapeutic agents for the treatment of immunological disease Browning, Jeffrey L., Brookline, MA, United States INVENTOR(S): Benjamin, Christopher D., Beverly, MA, United States Hochman, Paula S., Brookline, MA, United States Biogen, Inc., Cambridge, MA, United States (U.S. PATENT ASSIGNEE(S): corporation) NUMBER KIND DATE ______ US 5925351 19990720 US 1995-505606 19950721 (8) PATENT INFORMATION: APPLICATION INFO.: DOCUMENT TYPE: Utility Granted FILE SEGMENT: PRIMARY EXAMINER: Hutzell, Paula K.
ASSISTANT EXAMINER: Bansal, Geetha P.
LEGAL REPRESENTATIVE: Biogen, Inc., Flynn, Kerry A. 16 NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1,15 NUMBER OF DRAWINGS: 7 Drawing Figure(s); 7 Drawing Page(s) LINE COUNT:

ANSWER 41 OF 81 USPATFULL T.7

ACCESSION NUMBER: 1999:75310 USPATFULL

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Methods of treating TNF.alpha.-mediated disease using TITLE:

chimeric anti-TNF antibodies

INVENTOR(S):

Le, Junming, Jackson Heights, NY, United States Vilcek, Jan, New York, NY, United States Dadonna, Peter, Palo Alto, CA, United States Ghrayeb, John, Thorndale, PA, United States Knight, David, Berwyn, PA, United States Seigal, Scott, Westborough, MA, United States

New York University, New York, NY, United States (U.S. PATENT ASSIGNEE(S):

corporation)

Centocor, Inc., Malvern, PA, United States (U.S.

corporation)

NUMBER KIND DATE

US 5919452 19990706 US 1994-192861 19940204 (8) PATENT INFORMATION: APPLICATION INFO.:

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1993-10406, filed

on 29 Jan 1993, now abandoned And Ser. No. US

1993-13413, filed on 2 Feb 1993, now abandoned which is

a continuation-in-part of Ser. No. US 1992-943852, filed on 11 Sep 1992, now abandoned which is a

continuation-in-part of Ser. No. US 1992-853606, filed

on 18 Mar 1992, now abandoned which is a

continuation-in-part of Ser. No. US 1991-670827, filed

on 18 Mar 1991, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

Scheiner, Toni R. PRIMARY EXAMINER: Johnson, Nancy A. ASSISTANT EXAMINER:

Hamilton, Brook, Smith & Reynolds, P.C. LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: 13 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 48 Drawing Figure(s); 36 Drawing Page(s)

LINE COUNT: 5351

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 42 OF 81 USPATFULL

1999:69500 USPATFULL ACCESSION NUMBER:

Recombinant IL4 antibodies useful in treatment of IL4 TITLE:

mediated disorders

Holmes, Stephen D., Epsom, United Kingdom INVENTOR(S):

> Gross, Mitchell Stuart, Wayne, PA, United States Sylvester, Daniel R., Phoenixville, PA, United States

SmithKline Beecham Corporation, Philadelphia, PA, PATENT ASSIGNEE(S):

United States (U.S. corporation)

SmithKline Beecham p.l.c., Brentford, United Kingdom

(non-U.S. corporation)

NUMBER KIND

_____ US 5914110 US 1995-483636 19990622 PATENT INFORMATION: 19950607 (8) APPLICATION INFO.:

Continuation-in-part of Ser. No. WO 1994-US10308, filed RELATED APPLN. INFO.:

on 7 Sep 1994 which is a continuation-in-part of Ser. No. US 1993-136783, filed on 14 Oct 1993, now abandoned which is a continuation of Ser. No. US 1993-117366,

filed on 7 Sep 1993, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Caputa, Anthony C. Navarro, Mark ASSISTANT EXAMINER:

LEGAL REPRESENTATIVE: Eagle, Alissa M., King, William T., Venetianer, Stephen

NUMBER OF CLAIMS: 39 EXEMPLARY CLAIM: 1

12 Drawing Figure(s); 11 Drawing Page(s) NUMBER OF DRAWINGS:

2494 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 43 OF 81 USPATFULL

ACCESSION NUMBER: 1999:21735 USPATFULL

Method of treating viral diseases in animals TITLE: INVENTOR(S):

Whitfill, Craig E., Apex, NC, United States Thoma, John A., Fayetteville, AR, United States Fredericksen, Tommy L., Ashford, CT, United States Tyczkowski, Julius K., Cary, NC, United States

Thaxton, Jr., J. Paul, Brandon, MS, United States PATENT ASSIGNEE(S):

Embrex, Inc, Research Triangle Park, NC, United States

(U.S. corporation)

The University of Arkansas, Fayetteville, AR, United

States (U.S. corporation)

NUMBER KIND DATE -----

US 5871748 19990216 US 1996-697268 19960821 (8) PATENT INFORMATION: APPLICATION INFO.:

Continuation of Ser. No. US 1994-345291, filed on 28 RELATED APPLN. INFO.: Nov 1994, now abandoned which is a continuation of Ser. No. US 1993-8394, filed on 25 Jan 1993, now patented, Pat. No. US 5397569 which is a continuation of Ser. No.

> US 1990-586859, filed on 21 Sep 1990, now abandoned which is a continuation-in-part of Ser. No. US

1990-480678, filed on 15 Feb 1990, now abandoned which is a continuation-in-part of Ser. No. US 1989-416035,

filed on 2 Oct 1989, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Smith, Lynette F. ASSISTANT EXAMINER: Nelson, Brett L.

Myers Bigel Sibley & Sajovec, LLP LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: 64 EXEMPLARY CLAIM: 1

12 Drawing Figure(s); 7 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT: 1859

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 44 OF 81 USPATFULL

1999:18720 USPATFULL ACCESSION NUMBER:

Altered polypeptides with increased half-life TITLE:

Presta, Leonard G., San Francsico, CA, United States INVENTOR(S): Snedecor, Bradley R., Portola Valley, CA, United States

Genentech, Inc., So. San Francisco, CA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE

US 5869046 19990209 US 1995-422092 19950414 (8) PATENT INFORMATION: APPLICATION INFO.:

DOCUMENT TYPE: Utility Granted FILE SEGMENT:

PRIMARY EXAMINER: Eisenschenk, Frank C.
ASSISTANT EXAMINER: Rabin, Evelyn
LEGAL REPRESENTATIVE: Dreger, Walter H.

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

4 Drawing Figure(s); 3 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 45 OF 81 USPATFULL L7

1999:4367 USPATFULL ACCESSION NUMBER:

DNA encoding an insulin receptor substrate TITLE:

INVENTOR(S):

White, Morris F., West Roxbury, MA, United States Sun, Xiao Jian, Boston, MA, United States

Pierce, Jacalyn H., Potomac, MD, United States

Joslin Diabetes Center, Inc., Boston, MA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

KIND DATE NUMBER ----- PATENT INFORMATION: US 5858701 19990112 APPLICATION INFO.: US 1994-317310 19941003 (8) DOCUMENT TYPE: Utility

DOCUMENT TYPE:

Granted

FILE SEGMENT:

PRIMARY EXAMINER: Hutzell, Paula K. ASSISTANT EXAMINER: Hayes, Robert C.

LEGAL REPRESENTATIVE: Myers, Esq., Louis

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 8 Drawing Figure(s); 27 Drawing Page(s) LINE COUNT: 2713

INVENTOR(S):

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 46 OF 81 USPATFULL

ACCESSION NUMBER: 1998:159916 USPATFULL

TITLE:

Method of enhancing proliferation or differentiation of

hematopoietic stem cells using Wnt polypeptides Matthews, William, Woodside, CA, United States

Austin, Timothy W., Morgan Hill, CA, United States

Genentech, Inc., South San Francisco, CA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE ------

PATENT INFORMATION: US 5851984 19981222
APPLICATION INFO.: US 1996-696566 19960816 (8)
DOCUMENT TYPE: Utility

DOCUMENT TYPE: FILE SEGMENT: Granted

FILE SEGMENT: Granted
PRIMARY EXAMINER: Fitzgerald, David L.
ASSISTANT EXAMINER: Basham, Daryl A.
LEGAL REPRESENTATIVE: Svoboda, Craig G., Marschang, Diane L.
NUMBER OF CLAIMS: 20
EXEMPLARY CLAIM: 1

AVINGS: 4 Drawing Figure(s); 2 Drawing Page(s) 3923 NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 47 OF 81 USPATFULL

ACCESSION NUMBER: 1998:153858 USPATFULL

TITLE:

Antibodies to the antigen campath-1

Waldmann, Herman, Cambridge, United Kingdom INVENTOR(S): Clark, Michael R., Cambridge, United Kingdom Winter, Gregory P., Cambridge, United Kingdom

Riechmann, Lutz, La Jolla, CA, United States British Technology Group Limited, London, United

Kingdom (non-U.S. corporation)

NUMBER KIND DATE -----

PATENT ASSIGNEE(S):

PATENT INFORMATION: US 5846534 19981208 APPLICATION INFO.: US 1994-235705 19940429 (8)

RELATED APPLN. INFO.:

Continuation of Ser. No. US 1993-99480, filed on 30 Jul 1993, now abandoned which is a continuation of Ser. No. US 1992-921601, filed on 3 Aug 1992, now abandoned which is a continuation of Ser. No. US 1989-424233,

filed on 12 Oct 1989, now abandoned

NUMBER DATE -----GB 1988-3228 19880212 GB 1988-4464 19880225 PRIORITY INFORMATION:

DOCUMENT TYPE: FILE SEGMENT:

Utility Granted

PRIMARY EXAMINER: Budens, Robert D.

LEGAL REPRESENTATIVE: Nixon & Vanderhye

NUMBER OF CLAIMS: 18 EXEMPLARY CLAIM: 1,13

NUMBER OF DRAWINGS: 21 Drawing Figure(s); 13 Drawing Page(s)

1094 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 48 OF 81 USPATFULL

1998:150724 USPATFULL ACCESSION NUMBER: Chimeric antibodies TITLE:

Hardman, Norman, Riehen, Switzerland INVENTOR(S):

Gill, Laura Lee, Riehen, Switzerland

de Winter, Ronald F.J., Milton Ernest, England Wagner, Kathrin, Basel, Switzerland

Heusser, Christoph, Bottmingen, Switzerland

CIBA-GEIGY Corporation, Tarrytown, NY, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE _____ ___

US 5843708 19981201 US 1995-462371 19950605 (8) PATENT INFORMATION: APPLICATION INFO.:

Division of Ser. No. US 1994-307087, filed on 16 Sep RELATED APPLN. INFO.:

1994 which is a continuation of Ser. No. US

1992-947897, filed on 18 Sep 1992, now abandoned which is a continuation of Ser. No. US 1988-287178, filed on

21 Dec 1988, now abandoned

NUMBER DATE

GB 1988-77 19880105 GB 1988-20099 19880824 PRIORITY INFORMATION:

Utility

DOCUMENT TYPE: Granted FILE SEGMENT:

FILE SEGMENT: Granted
PRIMARY EXAMINER: Ziska, Suzanne E. LEGAL REPRESENTATIVE: Nowak, Henry P.

NUMBER OF CLAIMS: 21 EXEMPLARY CLAIM:

9 Drawing Figure(s); 7 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 49 OF 81 USPATFULL

ACCESSION NUMBER: 1998:150462 USPATFULL

Cellular and serum protein anchors for modulating TITLE:

pharmacokinetics

INVENTOR(S): Pouletty, Philippe, Atherton, CA, United States

Pouletty, Christine, Atherton, CA, United States RedCell Canada, Inc., Montreal, Canada (non-U.S.

PATENT ASSIGNEE(S):

corporation)

NUMBER KIND DATE -----US 5843440 US 1996-702127 19981201 PATENT INFORMATION:

19960814 (8) APPLICATION INFO.: Continuation of Ser. No. US 1993-137821, filed on 15 RELATED APPLN. INFO.:

Oct 1993, now abandoned which is a continuation-in-part of Ser. No. US 1993-70092, filed on 27 May 1993, now abandoned which is a continuation-in-part of Ser. No. US 1990-592214, filed on 3 Oct 1990, now abandoned

DOCUMENT TYPE: Utility Granted FILE SEGMENT:

PRIMARY EXAMINER: Scheiner, Toni R. Johnson, Nancy A. ASSISTANT EXAMINER:

LEGAL REPRESENTATIVE: Limbach & Limbach L.L.P.

NUMBER OF CLAIMS: 8
EXEMPLARY CLAIM: 1
LINE COUNT: 765

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 50 OF 81 USPATFULL

ACCESSION NUMBER: 1998:143883 USPATFULL

TITLE: Method of identifying modulators of binding between and

VCAM-1

INVENTOR(S): Gallatin, W. Michael, Mercer Island, WA, United States

Van der Vieren, Monica, Seattle, WA, United States

PATENT ASSIGNEE(S): ICOS Corporation, Bothell, WA, United States (U.S.

corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5837478 19981117 APPLICATION INFO.: US 1997-943363 19971003 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1996-605672, filed

on 22 Feb 1996 which is a continuation-in-part of Ser. No. US 1994-362652, filed on 21 Dec 1994, now patented, Pat. No. US 5766850 which is a continuation-in-part of Ser. No. US 1994-286889, filed on 5 Aug 1994, now patented, Pat. No. US 5470953, issued on 28 Nov 1995

which is a continuation-in-part of Ser. No. US 1993-173497, filed on 23 Dec 1993, now patented, Pat.

No. US 5437958, issued on 1 Aug 1995

DOCUMENT TYPE: Utility
FILE SEGMENT: Granted
PRIMARY EXAMINER: Feisee, Lila
ASSISTANT EXAMINER: Gambel, Phillip

LEGAL REPRESENTATIVE: Marshall, O'Toole, Gerstein, Murray & Borun

NUMBER OF CLAIMS: 4 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT: 7878

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 51 OF 81 USPATFULL

ACCESSION NUMBER: 1998:135170 USPATFULL

TITLE: Human .beta.2 integrin .alpha. subunit

INVENTOR(S): Gallatin, W. Michael, Mercer Island, WA, United States

Van der Vieren, Monica, Seattle, WA, United States ICOS Corporation, Bothell, WA, United States (U.S.

PATENT ASSIGNEE(S): ICOS Corpora corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5831029 19981103 APPLICATION INFO.: US 1995-482293 19950607 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1994-362652, filed on 21 Dec 1994 which is a continuation-in-part of Ser.

No. US 1994-286889, filed on 5 Aug 1994, now patented, Pat. No. US 5470953 which is a continuation-in-part of Ser. No. US 1993-173497, filed on 23 Dec 1993, now

patented, Pat. No. US 5437958

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Eisenschenk, Frank C.

ASSISTANT EXAMINER: Rabin, Evelyn

LEGAL REPRESENTATIVE: Marshall, O'Toole, Gerstein, Murray & Borun

NUMBER OF CLAIMS: 10 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT: 5481

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 52 OF 81 USPATFULL

ACCESSION NUMBER: 1998:122270 USPATFULL

TITLE: Human B2 integrin alpha subunit antibodies

INVENTOR(S): Gallatin, W. Michael, Mercer Island, WA, United States

Van der Vieren, Monica, Seattle, WA, United States

PATENT ASSIGNEE(S): ICOS Corporation, Bothell, WA, United States (U.S.

corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5817515 19981006
APPLICATION INFO.: US 1996-605672 19960222 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1994-362652, filed

on 21 Dec 1994 which is a continuation-in-part of Ser. No. US 1994-286889, filed on 5 Aug 1994, now patented, Pat. No. US 5470953, issued on 28 Nov 1995 which is a continuation-in-part of Ser. No. US 1993-173497, filed on 23 Dec 1993, now patented, Pat. No. US 5437958,

issued on 1 Aug 1995

DOCUMENT TYPE: Utility
FILE SEGMENT: Granted
PRIMARY EXAMINER: Feisee, Lila
ASSISTANT EXAMINER: Gambel, Phillip

LEGAL REPRESENTATIVE: Marshall, O'Toole, Gerstein, Murray & Borun

NUMBER OF CLAIMS: 2 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT: 6188

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 53 OF 81 USPATFULL

ACCESSION NUMBER: 1998:115416 USPATFULL

TITLE: Blockade of T lymphocyte down-regulation associated

with CTLA-4 signaling

INVENTOR(S): Allison, James Patrick, Berkeley, CA, United States

Leach, Dana R., Albany, CA, United States

Krummel, Matthew F., Berkeley, CA, United States

PATENT ASSIGNEE(S): The Regents of the University of California, Oakland,

CA, United States (U.S. corporation)

APPLICATION INFO.: US 1996-646605 19960508 (8)
RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1995-566853, filed

on 4 Dec 1995 which is a continuation-in-part of Ser. No. US 1995-506666, filed on 25 Jul 1995, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Scheiner, Toni R.

ASSISTANT EXAMINER: Lucas, John

LEGAL REPRESENTATIVE: Flehr Hohbach Test Albritton & Herbert, LLP

NUMBER OF CLAIMS: 11 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 27 Drawing Figure(s); 13 Drawing Page(s)

LINE COUNT: 1922

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 54 OF 81 USPATFULL

ACCESSION NUMBER: 1998:104391 USPATFULL

TITLE:

INVENTOR(S):

Antibodies to P-selectin and their uses

Chestnut, Robert W., Cardiff, CA, United States Polley, Margaret J., La Jolla, CA, United States Paulson, James C., Del Mar, CA, United States Jones, S. Tarran, Radlett, United Kingdom Saldanha, Jose W., Middlesex, United Kingdom

Bendig, Mary M., London, United Kingdom

Kriegler, Michael, Rancho Santa Fe, CA, United States

Perez, Carl, San Diego, CA, United States Bayer, Robert, San Diego, CA, United States Nunn, Michael, San Diego, CA, United States

Cytel Corporation, San Diego, CA, United States (U.S.

corporation)

NUMBER KIND DATE

PATENT INFORMATION: APPLICATION INFO.:

PATENT ASSIGNEE(S):

------US 5800815 19980901 US 1994-202047 19940225 (8)

Continuation-in-part of Ser. No. US 1993-57292, filed RELATED APPLN. INFO.:

on 5 May 1993, now abandoned which is a

continuation-in-part of Ser. No. US 1992-880198, filed

on 5 May 1992, now abandoned

NUMBER DATE

PRIORITY INFORMATION:

IL 1903-105614 19030505 WO 1993-US4274 19930504

DOCUMENT TYPE: FILE SEGMENT:

Utility Granted

PRIMARY EXAMINER: PRIMARY EXAMINER: Hutzell, Paula K ASSISTANT EXAMINER: Gambel, Phillip LEGAL REPRESENTATIVE: Campbell & Flore LEGAL REPRESENTATIVE:

Hutzell, Paula K. Campbell & Flores LLP

NUMBER OF CLAIMS:

57

EXEMPLARY CLAIM:

32,40 47 Drawing Figure(s); 40 Drawing Page(s)

NUMBER OF DRAWINGS: LINE COUNT:

4013

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 55 OF 81 USPATFULL

ACCESSION NUMBER:

1998:82874 USPATFULL

TITLE:

Monoclonal antibodies to cytotoxic lymphocyte

maturation factor

INVENTOR(S):

Gately, Maurice Kent, Montville, NJ, United States Gubler, Ulrich Andreas, Glen Ridge, NJ, United States Hulmes, Jeffrey David, Ringwood, NJ, United States Podlaski, Frank John, New City, NY, United States Stern, Alvin Seth, Passaic Park, NJ, United States Chizzonite, Richard Anthony, South Kent, CT, United

States

Pan, Yu-Ching Eugene, Pine Brook, NJ, United States Hoffmann-La Roche Inc., Nutley, NJ, United States (U.S.

corporation)

PATENT ASSIGNEE(S):

NUMBER KIND DATE _____ US 5780597 19980714 US 1995-460061 19950602 (8) PATENT INFORMATION:

APPLICATION INFO.: RELATED APPLN. INFO.:

Division of Ser. No. US 1994-205011, filed on 2 Mar 1994, now abandoned which is a division of Ser. No. US 1992-857023, filed on 24 Mar 1992, now abandoned which is a continuation-in-part of Ser. No. US 1990-572284, filed on 27 Aug 1990, now abandoned which is a

continuation-in-part of Ser. No. US 1990-520935, filed

on 9 May 1990, now abandoned which is a

continuation-in-part of Ser. No. US 1989-455708, filed

on 22 Dec 1989, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Cunningham, Thomas M. ASSISTANT EXAMINER: Lubet, Martha T.

LEGAL REPRESENTATIVE: Johnston, George W., Epstein, William H., Buchholz,

Briana C.

NUMBER OF CLAIMS: 3
EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 41 Drawing Figure(s); 44 Drawing Page(s)

LINE COUNT: 2912

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 56 OF 81 USPATFULL

ACCESSION NUMBER: 1998:82339 USPATFULL

TITLE: Methods of treatment of down syndrome by interferon

antagonists

INVENTOR(S): Maroun, Leonard E., Springfield, IL, United States PATENT ASSIGNEE(S): Meiogen Biotechnology Corporation, Springfield, IL,

United States (U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5780027 19980714 APPLICATION INFO.: US 1995-502519 19950714 (8)

DOCUMENT TYPE: Utility
FILE SEGMENT: Granted
PRIMARY EXAMINER: Huff, Sheela
ASSISTANT EXAMINER: Eyler, Yvonne

LEGAL REPRESENTATIVE: Pennie & Edmonds LLP

NUMBER OF CLAIMS: 16 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 777

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 57 OF 81 USPATFULL

ACCESSION NUMBER: 1998:68777 USPATFULL

TITLE: Human .beta.2 integrin .alpha. subunit

INVENTOR(S): Gallatin, W. Michael, Seattle, WA, United States

Van der Vieren, Monica, Seattle, WA, United States

PATENT ASSIGNEE(S): ICOS Corporation, Bothell, WA, United States (U.S.

corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5766850 19980616
APPLICATION INFO.: US 1994-362652 19941221 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1994-286889, filed on 5 Aug 1994, now patented, Pat. No. US 5470953 which is a continuation-in-part of Ser. No. US 1993-173497,

filed on 23 Dec 1993, now patented, Pat. No. US 5437958

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Furman, Keith C.

LEGAL REPRESENTATIVE: Marshall, O'Toole, Gerstein, Murray & Borun

NUMBER OF CLAIMS: 1 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT: 3283

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ACCESSION NUMBER:

1998:47965 USPATFULL

TITLE:

Polypeptides with increased half-life for use in treating disorders involving the LFA-1 receptor

INVENTOR(S):

Presta, Leonard G., San Francsico, CA, United States

Snedecor, Bradley R., Portola Valley, CA, United States Genentech, Inc., South San Francisco, CA, United States

(U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: APPLICATION INFO.:

PATENT ASSIGNEE(S):

US 5747035 19980505 US 1995-422091 19950414 (8)

DOCUMENT TYPE: FILE SEGMENT:

Utility Granted

PRIMARY EXAMINER: Feisee, Lila ASSISTANT EXAMINER: Gambel, Phillip LEGAL REPRESENTATIVE: Dreger, Walter H.

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

17 1

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 3 Drawing Page(s) LINE COUNT: 3305

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 59 OF 81 USPATFULL

ACCESSION NUMBER: 1998:39666 USPATFULL

TITLE:

Altered polypeptides with increased half-life Presta, Leonard G., San Francisco, CA, United States

INVENTOR(S): Snedecor, Bradley R., Portola Valley, CA, United States Genentech Inc., San Francisco, CA, United States (U.S. PATENT ASSIGNEE(S):

corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5739277 19980414
APPLICATION INFO.: US 1995-422101 19950414 (8)
DOCUMENT TYPE: Utility

DOCUMENT TYPE: FILE SEGMENT:

Granted

PRIMARY EXAMINER: Feisee, Lila
ASSISTANT EXAMINER: Johnson, Nancy A.
LEGAL REPRESENTATIVE: Hasak, Janet E.

NUMBER OF CLAIMS:

1,2,3

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

4 Drawing Figure(s); 3 Drawing Page(s)

LINE COUNT:

3251

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 60 OF 81 USPATFULL

ACCESSION NUMBER:

1998:27915 USPATFULL

TITLE:

Human .beta..sub.2 integrin .alpha.subunit

INVENTOR(S):

Gallatin, W. Michael, Mercer Island, WA, United States Van der Vieren, Monica, Seattle, WA, United States

PATENT ASSIGNEE(S):

ICOS Corporation, Bothell, WA, United States (U.S.

corporation)

NUMBER KIND DATE

PATENT INFORMATION:

_______ 19980317

APPLICATION INFO.:

US 5728533 US 1995-485618 19950607 (8)

RELATED APPLN. INFO.:

Continuation-in-part of Ser. No. US 1994-362652, filed on 21 Dec 1994 which is a continuation-in-part of Ser. No. US 1994-286889, filed on 5 Aug 1994, now patented, Pat. No. US 5470953 which is a continuation-in-part of Ser. No. US 1993-173497, filed on 23 Dec 1993, now

patented, Pat. No. US 5437958

DOCUMENT TYPE:

Utility Granted

FILE SEGMENT:

LEGAL REPRESENTATIVE: Marshall, O'Toole, Gerstein, Murray & Borun

PRIMARY EXAMINER: Carlson, Karen C.

NUMBER OF CLAIMS:

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

4 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT:

3915

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 61 OF 81 USPATFULL

ACCESSION NUMBER:

97:118165 USPATFULL

TITLE:

Product and process for targeting an immune response

INVENTOR(S):

Nemazee, David A., Denver, CO, United States

PATENT ASSIGNEE(S):

National Jewish Center for Immunology and Respiratory Medicine, Denver, CO, United States (U.S. corporation)

NUMBER KIND DATE ______

PATENT INFORMATION:
APPLICATION INFO.:

US 5698679 19971216 US 1994-309006 19940919 (8)

DOCUMENT TYPE: FILE SEGMENT:

Utility Granted

PRIMARY EXAMINER: Feisee, Lila
ASSISTANT EXAMINER: Eyler, Yvonne
LEGAL REPRESENTATIVE: Sheridan Ross P.C.

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

27

NUMBER OF DRAWINGS:

5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT:

1793

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 62 OF 81 USPATFULL

ACCESSION NUMBER:

97:117693 USPATFULL

TITLE:

Methods of treating rheumatoid arthritis using chimeric

anti-TNF antibodies

INVENTOR(S):

Le, Junming, Jackson Heights, NY, United States Vilcek, Jan, New York, NY, United States

Daddona, Peter, Menlo Park, CA, United States Ghrayeb, John, Thorndale, PA, United States Knight, David, Berwyn, PA, United States Siegel, Scott, Westborough, MA, United States

PATENT ASSIGNEE(S):

New York University Medical Center, New York, NY,

United States (U.S. corporation)

Centocor, Inc., Malvern, PA, United States (U.S.

corporation)

NUMBER KIND DATE _____

PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.: US 5698195 19971216 US 1994-324799 19941018 19941018 (8)

Continuation-in-part of Ser. No. US 1994-192102, filed on 4 Feb 1994 Ser. No. Ser. No. US 1994-192061, filed on 4 Feb 1994, now abandoned And Ser. No. US

1994-192093, filed on 4 Feb 1994, now abandoned , each Ser. No. US - which is a continuation-in-part of Ser. No. US 1993-10406, filed on 29 Jan 1993, now abandoned And Ser. No. US 1993-13413, filed on 2 Feb 1993, now abandoned which is a continuation-in-part of Ser. No. US 1992-943852, filed on 11 Sep 1992, now abandoned which is a continuation-in-part of Ser. No. US

1992-853606, filed on 18 Mar 1992, now abandoned which is a continuation-in-part of Ser. No. US 1991-670827,

filed on 18 Mar 1991, now abandoned

DOCUMENT TYPE: Utility Granted FILE SEGMENT: PRIMARY EXAMINER: Feisee, Lila ASSISTANT EXAMINER: Lucas, John

LEGAL REPRESENTATIVE: Hamilton, Brook, Smith & Reynolds, P.C. 16

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 33 Drawing Figure(s); 36 Drawing Page(s)

5887 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 63 OF 81 USPATFULL

97:114931 USPATFULL ACCESSION NUMBER:

Modified anti-ICAM-1 antibodies and their use in the TITLE:

 ${\tt treatment}\ {\tt of}\ {\tt inflammation}$

Faanes, Ronald Bertrand, Pound Ridge, NY, United States INVENTOR(S):

McGoff, Paul Edward, Watertown, CT, United States Shirley, Bret Allen, New Milford, CT, United States Scher, David Stuart, Danbury, CT, United States

PATENT ASSIGNEE(S): Boehringer Inglehiem Pharmaceuticals, Inc., Ridgefield,

CT, United States (U.S. corporation)

NUMBER KIND DATE ______

US 5695760 · 19971209 US 1995-427355 · 19950424 (8) PATENT INFORMATION:
APPLICATION INFO.: APPLICATION INFO.:

DOCUMENT TYPE: Utility FILE SEGMENT: Granted PRIMARY EXAMINER: Feisee, Lila ASSISTANT EXAMINER: Johnson, Nancy A.

Howrey & Simon, Auerbach, Jeffery I. LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS:

EXEMPLARY CLAIM: 1,12,24

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 3085

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 64 OF 81 USPATFULL

97:78179 USPATFULL ACCESSION NUMBER:

Monoclonal antibody compositions cross-reactive and TITLE:

cross-protective against P. aeruginosa serotypes

Siadak, Anthony W., Seattle, WA, United States INVENTOR(S):

Rosok, Mae J., Seattle, WA, United States Bristol-Myers Squibb Company, New York, NY, United PATENT ASSIGNEE(S):

States (U.S. corporation)

NUMBER KIND DATE -----US 5662905 US 1994-366204 19970902 PATENT INFORMATION: APPLICATION INFO.: 19941229 (8)

Continuation of Ser. No. US 1993-66604, filed on 24 May RELATED APPLN. INFO.:

1993, now patented, Pat. No. US 5378812 which is a continuation of Ser. No. US 1986-931179, filed on 24 Nov 1986, now abandoned which is a continuation-in-part of Ser. No. US 1985-807394, filed on 10 Dec 1985, now

abandoned Utility

FILE SEGMENT: Granted PRIMARY EXAMINER:

Loring, Susan A. LEGAL REPRESENTATIVE: Townsend And Townsend And Crew LLP

NUMBER OF CLAIMS: 8 EXEMPLARY CLAIM: LINE COUNT: 1412

DOCUMENT TYPE:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 65 OF 81 USPATFULL

ACCESSION NUMBER: 97:70718 USPATFULL

TITLE: Methods of treating TNF-.alpha.-mediated Crohn's

disease using chimeric anti-TNF antibodies

INVENTOR(S): Le, Junming, Jackson Heights, NY, United States Vilcek, Jan, New York, NY, United States

Vilcek, Jan, New York, NY, United States Dadonna, Peter, Palo Alto, CA, United States Ghrayeb, John, Thorndale, PA, United States Knight, David, Berwyn, PA, United States

Siegel, Scott A., Westborough, MA, United States (S): New York University Medical Center, New York, NY,

PATENT ASSIGNEE(S): New York University Medical Center, New United States (U.S. corporation)

Centocor, Inc., Malvern, PA, United States (U.S.

corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5656272 19970812
APPLICATION INFO.: US 1994-192102 19940204 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1993-10406, filed

on 26 Jan 1993, now abandoned And Ser. No. US

1993-13413, filed on 2 Feb 1993, now abandoned which is

a continuation-in-part of Ser. No. US 1992-943852,

filed on 11 Sep 1992, now abandoned which is a

continuation-in-part of Ser. No. US 1992-853606, filed

on 18 Mar 1992, now abandoned which is a

continuation-in-part of Ser. No. US 1991-670827, filed

on 18 Mar 1991, now abandoned

DOCUMENT TYPE: FILE SEGMENT:

FILE SEGMENT: Granted
PRIMARY EXAMINER: Feisee, Lila
ASSISTANT EXAMINER: Lucas, John

LEGAL REPRESENTATIVE: Hamilton, Brook, Smith & Reynolds, P.C.

Utility

NUMBER OF CLAIMS: 7 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 48 Drawing Figure(s); 36 Drawing Page(s)

LINE COUNT: 5251

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 66 OF 81 USPATFULL

ACCESSION NUMBER: 97:40481 USPATFULL

TITLE: Method for inhibiting the viability of Pseudomonas

aeruginosa with cross-reactive and cross-protective

monoclonal antibodies

INVENTOR(S): Siadak, Anthony W., Seattle, WA, United States

Rosok, Mae J., Seattle, WA, United States

PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, New York, NY, United

States (U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: APPLICATION INFO.:

US 5628996 19970513 US 1995-463910 19950605 (8)

RELATED APPLN. INFO.:

Division of Ser. No. US 1994-366204, filed on 29 Dec 1994 which is a continuation of Ser. No. US 1993-66604, filed on 24 May 1993, now patented, Pat. No. US 5378812

which is a continuation of Ser. No. US 1986-931179, filed on 24 Nov 1986, now abandoned which is a

continuation-in-part of Ser. No. US 1985-807394, filed

on 10 Dec 1985, now abandoned

DOCUMENT TYPE: FILE SEGMENT:

Utility Granted

PRIMARY EXAMINER:

Housel, James C.

ASSISTANT EXAMINER:

Loring, Susan A.

LEGAL REPRESENTATIVE:

Townsend and Townsend and Crew LLP

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

10 1

1405

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 67 OF 81 USPATFULL

ACCESSION NUMBER:

97:38410 USPATFULL

TITLE:

Monoclonal antibodies cross-reactive and

cross-protective against human monoclonal antibodies

against pseudomonas aeruginosa serotypes

INVENTOR(S):

Siadak, Anthony W., Seattle, WA, United States

Rosok, Mae J., Seattle, WA, United States

PATENT ASSIGNEE(S):

Bristol-Myers Squibb Company, New York, NY, United

States (U.S. corporation)

NUMBER KIND _____ -----

PATENT INFORMATION:

US 5627067 19970506

APPLICATION INFO .:

US 5627067 19970506 US 1995-462370 19950605 (8)

RELATED APPLN. INFO.:

Division of Ser. No. US 1994-366204, filed on 29 Dec 1994 which is a continuation of Ser. No. US 1993-66604, filed on 24 May 1993, now patented, Pat. No. US 5378812

which is a continuation of Ser. No. US 1986-931179,

filed on 24 Nov 1986, now abandoned which is a

continuation-in-part of Ser. No. US 1985-807394, filed

on 10 Dec 1985, now abandoned

DOCUMENT TYPE:

Utility Granted

FILE SEGMENT:

Loring, Susan A.

PRIMARY EXAMINER: LEGAL REPRESENTATIVE:

Townsend and Townsend and Crew

NUMBER OF CLAIMS:

EXEMPLARY CLAIM: LINE COUNT:

1391

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 68 OF 81 USPATFULL T.7

ACCESSION NUMBER:

97:31409 USPATFULL

TITLE:

Inhibition of intimal hyperplasia using antibodies to

PDGF beta receptors

INVENTOR(S):

Hart, Charles E., Brier, WA, United States Kenagy, Richard D., Seattle, WA, United States

PATENT ASSIGNEE(S):

Clowes, Alexander W., Seattle, WA, United States ZymoGenetics, Inc., Seattle, WA, United States (U.S.

corporation)

University of Washington, Seattle, WA, United States

(U.S. corporation)

NUMBER KIND DATE _____ 19970415

PATENT INFORMATION:

US 5620687 US 1994-366860 19941230 (8)

APPLICATION INFO.: RELATED APPLN. INFO.:

Continuation-in-part of Ser. No. US 1994-304623, filed on 12 Sep 1994, now abandoned which is a continuation of Ser. No. US 1993-23504, filed on 25 Feb 1993, now

abandoned

DOCUMENT TYPE: FILE SEGMENT:

Utility Granted

PRIMARY EXAMINER: ASSISTANT EXAMINER: Chan, Christina Y.

LEGAL REPRESENTATIVE:

Gambel, Phillip Parker, Gary E., Leith, Debra K., Sawislak, Deborah A.

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

19 1

12 Drawing Figure(s); 12 Drawing Page(s) NUMBER OF DRAWINGS:

2786 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 69 OF 81 USPATFULL

97:6049 USPATFULL ACCESSION NUMBER:

TITLE:

Method of refolding human IL-13

INVENTOR(S):

Culpepper, Janice, Mountain View, CA, United States

McKenzie, Andrew, Redwood City, CA, United States

Dang, Warren, San Jose, CA, United States

Zurawski, Gerard, Redwood City, CA, United States

Schering Corporation, Kenilworth, NJ, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE _____ 19970121

PATENT INFORMATION: APPLICATION INFO.:

US 5596072 US 1993-12543 19930201 (8)

RELATED APPLN. INFO.:

Continuation-in-part of Ser. No. US 1992-933416, filed

on 21 Aug 1992, now abandoned

DOCUMENT TYPE:

Utility Granted

ASSISTANT EXAMINER: Draper, Garnette D. Spector Towns. LEGAL REPORTS.

Spector, Lorraine M.

LEGAL REPRESENTATIVE: Ching, Edwin P.

NUMBER OF CLAIMS:

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

288 Drawing Figure(s); 61 Drawing Page(s)

LINE COUNT:

4619

10

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 70 OF 81 USPATFULL L7

ACCESSION NUMBER:

97:3527 USPATFULL

TITLE:

Method of producing an anti-D immunoglobulin

concentrate and a pharmaceutical

preparation

INVENTOR(S):

Hodler, Gerhard, Worb, Switzerland Lerch, Peter, Bern, Switzerland

Stucki, Martin, Laupen, Switzerland

PATENT ASSIGNEE(S):

Rotkreuzstiftung Zentrallaboratorium Blutspendedienst,

Bern, Switzerland (non-U.S. corporation)

NUMBER KIND DATE US 5593675 US 1994-360334 PATENT INFORMATION: 19970114 APPLICATION INFO.: 19941221 (8)

> NUMBER DATE --**--**-----

PRIORITY INFORMATION: CH 1993-93810912 19931227 Utility

DOCUMENT TYPE:

FILE SEGMENT: Granted
PRIMARY EXAMINER: Feisee, Lila

LEGAL REPRESENTATIVE:

Seed and Berry LLP

NUMBER OF CLAIMS:

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

1 Drawing Figure(s); 1 Drawing Page(s)

LINE COUNT: 552

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 71 OF 81 USPATFULL

ACCESSION NUMBER:

96:82450 USPATFULL

TITLE:

Methods and vaccines comprising surface-active

copolymers

INVENTOR(S):

PATENT ASSIGNEE(S):

Hunter, Robert L., Tucker, GA, United States

Emory University, Atlanta, GA, United States (U.S.

corporation)

NUMBER KIND DATE

PATENT INFORMATION:

APPLICATION INFO.:

US 5554372 19960910 US 1995-420333 19950411 (8)

RELATED APPLN. INFO.:

Continuation of Ser. No. US 1993-133760, filed on 7 Oct 1993, now abandoned which is a continuation of Ser. No. US 1991-716807, filed on 21 Jun 1991, now abandoned

which is a continuation-in-part of Ser. No. US

1990-544831, filed on 27 Jun 1990, now abandoned which is a continuation-in-part of Ser. No. US 1989-449086,

filed on 8 Dec 1989, now abandoned which is a continuation of Ser. No. US 1989-341315, filed on 21 Apr 1989, now abandoned which is a continuation of Ser. No. US 1988-208335, filed on 17 Jun 1988, now abandoned

which is a continuation-in-part of Ser. No. US

1987-75187, filed on 16 Jul 1987, now abandoned which is a continuation-in-part of Ser. No. US 1986-909964,

filed on 22 Sep 1986, now abandoned

DOCUMENT TYPE:

FILE SEGMENT:

Utility Granted

2669

PRIMARY EXAMINER: ASSISTANT EXAMINER: Housel, James C. Shaver, Jennifer

LEGAL REPRESENTATIVE:

Jones & Askew

NUMBER OF CLAIMS:

NUMBER OF DRAWINGS:

EXEMPLARY CLAIM: 1

24 Drawing Figure(s); 17 Drawing Page(s)

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 72 OF 81 USPATFULL 1.7

ACCESSION NUMBER:

95:105943 USPATFULL

TITLE:

Human .beta..sub.2 integrin .alpha. subunit Gallatin, W. Michael, Seattle, WA, United States

INVENTOR(S):

PATENT ASSIGNEE(S):

Van der Vieren, Monica, Seattle, WA, United States ICOS Corporation, Bothell, WA, United States (U.S.

corporation)

NUMBER KIND DATE ______

PATENT INFORMATION: APPLICATION INFO.:

US 5470953 19951128 US 1994-286889 19940805 (8)

RELATED APPLN. INFO.:

Continuation-in-part of Ser. No. US 1993-173497, filed

on 23 Dec 1993

DOCUMENT TYPE: FILE SEGMENT:

Utility Granted

PRIMARY EXAMINER:

Wax, Robert A.

ASSISTANT EXAMINER:

Kim, Hyosuk Marshall, O'Toole, Gerstein, Murray & Borun

LEGAL REPRESENTATIVE: NUMBER OF CLAIMS:

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

4 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT:

2422

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 73 OF 81 USPATFULL L7

ACCESSION NUMBER:

95:22692 USPATFULL

TITLE:

Method of treating Infectious Bursal Disease Virus

infections

INVENTOR(S):

Whitfill, Craig E., 1300 Wellstone Cir., Apex, NC,

United States 27502

Thomas, John A., 1206 Crestwood Dr., Fayetteville, AR,

United States 72701

Fredericksen, Tommy L., 591 Westford Rd., Ashford, CT,

United States 06278

Tyczkowski, Julius K., 111 Woodruff Ct., Cary, NC,

United States 27511

Thaxton, Jr., J. Paul, 117 Campfire Cir., Brandon, MS,

United States 39240

NUMBER KIND DATE

US 5397569 PATENT INFORMATION: 19950314 US 1993-8394 APPLICATION INFO.: 19930125 (8)

Continuation of Ser. No. US 1990-586859, filed on 21 RELATED APPLN. INFO.: Sep 1990 which is a continuation-in-part of Ser. No. US

1990-480678, filed on 15 Feb 1990, now abandoned which is a continuation-in-part of Ser. No. US 1989-416035,

filed on 2 Oct 1989, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

Nucker, Christine M. PRIMARY EXAMINER: Krsek-Staples, Julie ASSISTANT EXAMINER:

Bell, Seltzer, Park & Gibson LEGAL REPRESENTATIVE:

49 NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 12 Drawing Figure(s); 5 Drawing Page(s)

1785 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 74 OF 81 USPATFULL

95:22691 USPATFULL ACCESSION NUMBER:

Method of treating infectious bursal disease virus TITLE:

infections

Whitfill, Craig E., 1300 Wellstone Cir., Apex, NC, INVENTOR(S):

United States 27502

Thoma, John A., 1206 Crestwood Dr., Fayetteville, AR, United States 72701

Fredericksen, Tommy L., 591 Westford Rd., Ashford, CT,

United States 06278

Tyczkowski, Julius K., 111 Woodruff Ct., Cary, NC,

United States 27511

Thaxton, Jr., J. Paul, 117 Campfire Cir., Brandon, MS,

United States 39240

NUMBER KIND DATE

US 5397568 US 1993-24093 PATENT INFORMATION: 19950314 19930225 (8) APPLICATION INFO.:

Continuation of Ser. No. US 1990-591523, filed on 1 Oct RELATED APPLN. INFO.:

1990 which is a continuation-in-part of Ser. No. US 1990-480678, filed on 15 Feb 1990, now abandoned which is a continuation-in-part of Ser. No. US 1989-416035,

filed on 2 Oct 1989, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

Nucker, Christine M. PRIMARY EXAMINER: Krsek-Staples, Julie Bell, Seltzer, Park & Gibson ASSISTANT EXAMINER:

LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: 34 EXEMPLARY CLAIM:

12 Drawing Figure(s); 5 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT: 1787

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 75 OF 81 USPATFULL

95:1714 USPATFULL ACCESSION NUMBER:

TITLE: Monoclonal antibodies cross-reactive and

cross-protective against P. aeruginosa serotypes Siadak, Anthony W., Seattle, WA, United States INVENTOR(S):

Rosok, Mae J., Seattle, WA, United States

Bristol-Myers Squibb Company, New York, NY, United PATENT ASSIGNEE(S):

States (U.S. corporation)

NUMBER _____

US 5378812 19950103 US 1993-66604 19930524 (8) PATENT INFORMATION: APPLICATION INFO .:

Continuation of Ser. No. US 1986-931179, filed on 24 RELATED APPLN. INFO .: Nov 1986, now abandoned which is a continuation-in-part

of Ser. No. US 1985-807391, filed on 10 Dec 1985, now

abandoned DOCUMENT TYPE: Utility Granted FILE SEGMENT:

PRIMARY EXAMINER: Lacey, David L. ASSISTANT EXAMINER: Loring, Susan A.

Townsend and Townsend Khourie and Crew LEGAL REPRESENTATIVE:

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1 LINE COUNT: 1363

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 76 OF 81 USPATFULL L7

94:33305 USPATFULL ACCESSION NUMBER:

DNA sequence encoding a selectin ligand TITLE:

Lasky, Laurence A., Sausalito, CA, United States Imai, Yasuyuki, San Francisco, CA, United States INVENTOR(S):

Rosen, Steven D., San Francisco, CA, United States

Singer, Mark S., Berkeley, CA, United States

Genentech, Inc., So. San Francisco, CA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

Regents of the University of California, Alameda, CA,

United States (U.S. corporation)

NUMBER KIND ______ US 5304640 US 1992-834902 PATENT INFORMATION: APPLICATION INFO.: 19940419

19920213 (7)

Continuation-in-part of Ser. No. US 1991-695805, filed RELATED APPLN. INFO.:

on 6 May 1991

DOCUMENT TYPE: Utility Granted FILE SEGMENT:

PRIMARY EXAMINER: Schwartz, Richard A. ASSISTANT EXAMINER: Guzo, David LEGAL REPRESENTATIVE: Dreger, Ginger R.

NUMBER OF CLAIMS: 20 EXEMPLARY CLAIM:

17 Drawing Figure(s); 12 Drawing Page(s) NUMBER OF DRAWINGS:

2371 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 77 OF 81 USPATFULL L7

ACCESSION NUMBER: 93:54852 USPATFULL

TITLE: Lymphocyte homing receptor/immunoglobulin fusion

proteins

Capon, Daniel J., San Mateo, CA, United States INVENTOR(S):

Lasky, Laurence A., Sausalito, CA, United States

PATENT ASSIGNEE(S): Genentech, Inc., South San Francisco, CA, United States

(U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5225538 19930706 APPLICATION INFO.: US 1991-808122 19911216 (7)

RELATED APPLN. INFO.: Division of Ser. No. US 1989-440625, filed on 22 Nov 1989, now patented, Pat. No. US 5116964 which is a continuation of Ser. No. US 1989-315015, filed on 23

Feb 1989, now patented, Pat. No. US 5098833

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Hill, Jr., Robert J.

ASSISTANT EXAMINER: Ulm, John D. LEGAL REPRESENTATIVE: Dreger, Ginger R.

NUMBER OF CLAIMS: 29 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 21 Drawing Figure(s); 18 Drawing Page(s)

LINE COUNT: 2558

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 78 OF 81 USPATFULL

ACCESSION NUMBER: 92:86949 USPATFULL

TITLE: Serine protease inhibitors

INVENTOR(S): Glover, George I., Creve Coeur, MO, United States

Schasteen, Charles S., University City, MO, United

States

PATENT ASSIGNEE(S): Monsanto Company, St. Louis, MO, United States (U.S.

corporation)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1988-200821, filed on 1 Jun

1988, now abandoned which is a continuation of Ser. No. US 1987-6725, filed on 6 Feb 1987, now abandoned which is a continuation-in-part of Ser. No. US 1986-840810,

filed on 18 Mar 1986, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Chan, Y. Christina LEGAL REPRESENTATIVE: Bennett, Dennis A.

NUMBER OF CLAIMS: 25 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 3 Drawing Figure(s); 3 Drawing Page(s)

LINE COUNT: 1966

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 79 OF 81 USPATFULL

ACCESSION NUMBER: 92:42890 USPATFULL
TITLE: Hybrid immunoglobulins

INVENTOR(S): Capon, Daniel J., San Mateo, CA, United States
Lasky, Laurence A., Sausalito, CA, United States

PATENT ASSIGNEE(S): Genentech, Inc., South San Francisco, CA, United States

(U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5116964 19920526 APPLICATION INFO.: US 1989-440625 19891122 (7)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1989-315015, filed

on 23 Feb 1989

DOCUMENT TYPE: Utility
FILE SEGMENT: Granted

ASSISTANT EXAMINER: Lacey, David L. Ulm, John D.

LEGAL REPRESENTATIVE: Dreger, Ginger R., Adler, Carolyn R.

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

• 679

14 Drawing Figure(s); 18 Drawing Page(s)
2533 NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 80 OF 81 USPATFULL

ACCESSION NUMBER: 92:10801 USPATFULL

Human monoclonal antibody to lymphadenopathy-associated TITLE:

virus

McClure, Janela, Vashon Island, WA, United States INVENTOR(S):

Genetic Systems Corporation, Redmond, WA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE ______

PATENT INFORMATION:
APPLICATION INFO.:
RELATED APPLN. INFO.: US 5087557 19920211 US 1990-498454 19900319 (7)

Continuation of Ser. No. US 1986-877579, filed on 23

Jun 1986, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Granted

PRIMARY EXAMINER: Nucker, Christine

LEGAL REPRESENTATIVE: Townsend and Townsend

NUMBER OF CLAIMS: 18 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 2 Drawing Figure(s); 2 Drawing Page(s) LINE COUNT: 631

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 81 OF 81 USPATFULL

88:2780 USPATFULL ACCESSION NUMBER:

Immunomodulating medication based on Fc fragments of TITLE:

human IqG

Carosella, Edgardo D., Lyons, France INVENTOR(S):

Armand, Jacques B., St Germain, France

Institut Merieux, Lyons, France (non-U.S. corporation) PATENT ASSIGNEE(S):

NUMBER KIND DATE ______ PATENT INFORMATION: US 4719107 APPLICATION INFO.: US 1984-679445 19880112

19841207 (6)

NUMBER DATE _____ ___

FR 1983-19568 19831207 PRIORITY INFORMATION:

DOCUMENT TYPE: Utility Granted

FILE SEGMENT: Granteo
PRIMARY EXAMINER: Schain, Howard E.
LEGAL REPRESENTATIVE: Cushman, Darby & Cushman

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 945 LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

FILE 'BIOSIS, CABA, CAPLUS, EMBASE, LIFESCI, MEDLINE, SCISEARCH,
USPATFULL, JAPIO' ENTERED AT 14:47:09 ON 25 OCT 2001

L1 5521 S CONCENTR? AND LYOPHILIZ? AND IMMUNOGLOBULIN?

L2 3813 S L1 AND IGG

L3 3756 S L2 AND PREPAR?

L4 708 S L3 AND (ION EXCHANGE CHROMATOGRAPHY)

L5 706 DUP REM L4 (2 DUPLICATES REMOVED)

L6 24 S L5 AND IGG4

=>

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ANSWER 1 OF 24 USPATFULL
L6
       2001:184842 USPATFULL
ΑN
TΙ
       Fas antigen derivatives
       Nakamura, Norio, Tokyo, Japan
IN
       Nagata, Shigekazu, Osaka-fu, Japan
       Mochida Pharmaceutical Co., Ltd., Tokyo, Japan (non-U.S. corporation)
PA
       Osaka Bioscience Institute, Osaka, Japan (non-U.S. corporation)
                               20011023
PΙ
       US 6306395
                          B1
       WO 9742319 19971113
       US 1998-180100
                               19981102 (9)
ΑI
       WO 1997-JP1502
                               19970501
                                         PCT 371 date
                               19981102
                               19981102 PCT 102(e) date
       JP 1996-135760
                           19960502
PRAI
DT
       Utility
FS
       GRANTED
       Primary Examiner: Huff, Sheela; Assistant Examiner: Harris, Alana M.
EXNAM
LREP
       Birch, Stewart, Kolasch & Birch, LLP
CLMN
       Number of Claims: 22
ECL
       Exemplary Claim: 1
DRWN
       15 Drawing Figure(s); 28 Drawing Page(s)
LN.CNT 2004
ΑB
       This invention provides a novel Fas antigen derivative which comprises
       at least a part or entire portion of Fas antigen extracellular region
       polypeptide in which at least one amino acid residue is deleted from a
       group of amino acid residues starting from the N-terminal amino acid
       residue of the Fas antigen polypeptide to a cysteine residue most close
       to the N-terminal side (excluding said cysteine residue), as well as a
       DNA fragment which encodes Fas antigen derivative, a recombinant DNA
       molecule which contains DNA sequence, a transformant in which
       recombinant DNA molecule is introduced, a method for the production of
       Fas antigen derivative, a medicament which contains novel Fas antigen
       derivative as the active ingredient and a method for the improvement of
       activities and functions of Fas antigen and the like.
L6
     ANSWER 2 OF 24 USPATFULL
       2001:165585 USPATFULL
AN
       Immunoassay technique using multispecific molecules
TΙ
       Khaw, Ban-an, Milton, MA, United States
IN
       Narula, Jagat, Rosemont, PA, United States
PΤ
       US 2001024795
                          Α1
                               20010927
                          Α1
                               20001201 (9)
ΑI
       US 2000-727421
       Continuation-in-part of Ser. No. US 1999-380168, filed on 6 Oct 1999,
RLI
       PENDING
                           19970226 (60)
PRAI
       US 1997-39111
DT
       Utility
FS
       APPLICATION
LREP
       PENNIE AND EDMONDS, 1155 AVENUE OF THE AMERICAS, NEW YORK, NY, 100362711
CLMN
       Number of Claims: 55
ECL
       Exemplary Claim: 1
DRWN
       3 Drawing Page(s)
LN.CNT 1620
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention relates to compositions and methods for detecting
       very low concentrations of a molecule in a mixture. The
       detection of the molecule comprises the steps of first contacting a
       sample with a multispecific molecule capable of binding at least two
       molecules including the molecule to be detected, wherein the molecule to
       be detected is bound by the multispecific molecule thereby forming a
       complex, and second contacting the complex with a second, different
       molecule which is linked via a polymer to multiple detection signaling
       molecules. The invention may also be practiced by administration of the
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multispecific molecule in vivo, to a host for the molecule to be

detected, either with or without the bound polymer probe and thereafter, respectively, either detecting the signaling molecule on the probe, or administering the probe and allowing it to bind the multispecific molecule, followed by detection of the signaling molecule on the probe.

```
ANSWER 3 OF 24 USPATFULL
L6
       2001:160973 USPATFULL
ΑN
TΤ
       Use of heregulin as a growth factor
ΙN
       Sliwkowski, Mark X., San Carlos, CA, United States
       Kern, Jeffrey A., Iowa City, IA, United States
                               20010920
PΙ
       US 2001023241
                          Α1
       US 2001-773517
                               20010202 (9)
ΑI
                          Α1
       Continuation of Ser. No. US 1999-243198, filed on 2 Feb 1999, ABANDONED
RLI
PRAI
       US 1998-73866
                           19980204 (60)
DT
       Utility
FS
       APPLICATION
       Supervisor, Patent Prosecution Services, PIPER MARBURY RUDNICK & WOLFE
LREP
       LLP, 1200 Nineteenth Street, N.W., Washington, DC, 20036-2412
CLMN
       Number of Claims: 20
ECL
       Exemplary Claim: 1
DRWN
       28 Drawing Page(s)
LN.CNT 3786
       Ligands which bind to the HER2, HER3 and/or HER4 receptors are useful as
AB
       normal epithelial cell growth factors.
L6
     ANSWER 4 OF 24 USPATFULL
       2001:142469 USPATFULL
ΑN
       Process for producing immunoglobulins for intravenous
ΤT
       administration and other immunoglobulin products
       Laursen, Inga, Hellerup, Denmark
IN
       Teisner, B.o slashed.rge, Odense C, Denmark
       Statens Serum Institut, Copenhagen S., Denmark (non-U.S. corporation)
PA
PΙ
       US 6281336
                               20010828
       US 1999-328497
                               19990609 (9)
ΑI
       EP 1998-201909
                           19980609
PRAI
       US 1998-102055
                           19980928 (60)
DT
       Utility
FS
       GRANTED
       Primary Examiner: Saunders, David
EXNAM
       Birch, Stewart, Kolasch & Birch, LLP
LREP
CLMN
       Number of Claims: 14
ECL
       Exemplary Claim: 1
       No Drawings
DRWN
LN.CNT 1465
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention relates to a process for purifying
AΒ
       immunoglobulin G from a crude immunoglobulin
       -containing plasma protein fraction. Said process includes a number of
       steps of which the anion exchange chromatography and the cation exchange
       chromatography are preferably connected in series. An acetate buffer
       having a pH of about 5.0-6.0 and having a molarity of about 5-25 mM is
       preferably used throughout the purification process. The invention
       further comprises an immunoglobulin product which is
       obtainable by this process. The invention also relates to an
       immunoglobulin product which has a purity of more than 98%, has
       a content of IgG monomers and dimers of more than 98.5%, has a
       content of IgA less than 4 mg of IgA/1, and contains less than 0.5%
       polymers and aggregates. Said product does not comprise detergent, PEG
       or albumin as a stabilizer. The product is stable, virus-safe, liquid
       and ready for instant intravenous administration.
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ANSWER 5 OF 24 USPATFULL L6 ΑN

g #

2001:141881 USPATFULL

ΤI Methods of using an AL-1 neurotrophic factor immunoadhesin

```
Caras, Ingrid W., San Francisco, CA, United States
ΙN
       Winslow, John W., El Granada, CA, United States
       Genentech, Inc., South San Francisco, CA, United States (U.S.
PA
       corporation)
                                20010828
PΙ
       US 6280732
                           В1
       US 1995-486449
                                19950607 (8)
AΙ
       Continuation-in-part of Ser. No. US 1994-330128, filed on 27 Oct 1994,
RLI
       now abandoned
DT
       Utility
FS
       GRANTED
       Primary Examiner: Caputa, Anthony C; Assistant Examiner: Gucker, Stephen
EXNAM
       Torchia, Timothy E.
LREP
CLMN
       Number of Claims: 5
ECL
       Exemplary Claim: 1
       7 Drawing Figure(s); 7 Drawing Page(s)
DRWN
LN.CNT 2167
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention provides nucleic acids encoding AL-1 protein, as
       well as AL-1 protein produced by recombinant DNA methods. Such AL-1
       protein is useful in preparing antibodies and in diagnosing
       and treating various neuronal disorders.
L6
     ANSWER 6 OF 24 USPATFULL
AN
       2001:59631 USPATFULL
TΙ
       Methods for identifying erythropoietin receptor binding protein
       Middleton, Steven A., Flemington, NJ, United States Johnson, Dana, Upper Black Eddy, PA, United States
TN
       McMahon, Frank J., Whitehouse Station, NJ, United States
       Mulkahy, Linda S., Yardley, PA, United States
Jolliffe, Linda K., Belle Mead, NJ, United States
       Ortho Pharmaceutical Corporation, Raritan, NJ, United States (U.S.
PA
       corporation)
                                 20010424
       US 6221608
                           В1
PΤ
       US 1997-786690
                                 19970122 (8)
ΑI
       Utility
DT
FS
       Granted
       Primary Examiner: Spector, Lorraine; Assistant Examiner: Kaufman, Claire
EXNAM
       Wallen, III, John W.
LREP
CLMN
       Number of Claims: 1
ECL
       Exemplary Claim: 1
       19 Drawing Figure(s); 12 Drawing Page(s)
DRWN
LN.CNT 1671
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The extracellular domain of the human erythropoietin receptor (EPO
AΒ
       binding protein, EBP) has been expressed and overproduced in E. coli.
       Control of oxygen levels and pH during high density fermentation allows
       the production of only the protein variant with the native amino
       terminus. Methods disclosed permit the efficient recovery of purified
       EBP which quantitatively binds EPO. The active purified protein competes
       with membrane associated EPO receptor for binding [.sup.125 I]EPO and
       neutralizes EPO dependent stimulation in a cell based proliferation
       assay. Further, the radioligand equilibrium binding constant for this
       interaction has been determined by immobilizing EBP on agarose gel via a
       free cysteine. The EBP of the present invention has many uses including
       the structural determination of the protein by NMR or crystallography,
       in drug design and discovery, and as a therapeutic. A fusion protein of
       EBP and an immunoglobulin heavy chain was also produced. This
       protein, termed EBP-Ig, is a preformed dimerization template and is also
       useful in drug design and discovery methods.
L6
     ANSWER 7 OF 24 USPATFULL
       2000:167510 USPATFULL
ΑN
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, •

TΙ

Uses of Wnt polypeptides

1

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Matthews, William, Woodside, CA, United States
IN
       Austin, Timothy W., Morgan Hill, CA, United States
       Genentech, Inc., So. San Francisco, CA, United States (U.S. corporation)
PA
                               20001212
PΙ
       US 6159462
                               19970815 (8)
       US 1997-911860
ΑI
       US 1996-24068
                           19960816 (60)
PRAI
DT
       Utility
FS
       Granted
       Primary Examiner: Saunders, David; Assistant Examiner: VanderVegt, F.
EXNAM
LREP
       Svoboda, Craig G., Carpenter, David A.
       Number of Claims: 13
CLMN
       Exemplary Claim: 1
ECL
       4 Drawing Figure(s); 2 Drawing Page(s)
DRWN
LN.CNT 3907
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Uses for Wnt polypeptides in hematopoiesis are disclosed. In particular,
       in vitro and in vivo methods for enhancing proliferation,
       differentiation or maintenance of a hematopoietic stem/progenitor cell
       using a Wnt polypeptide, and optionally another cytokine, are described.
     ANSWER 8 OF 24 USPATFULL
L6
ΑN
       2000:157557 USPATFULL
TΙ
       Monoclonal antibodies specific for the extracellular domain of
       prostate-specific membrane antigen
       Murphy, Gerald P., Seattle, WA, United States
TN
       Boynton, Alton L., Redmond, WA, United States
       Holmes, Eric H., Bothell, WA, United States
       Tino, William Thomas, Redmond, WA, United States
       Northwest Biotherapeutics, Inc., Seattle, WA, United States (U.S.
PA
       corporation)
                                20001121
PΙ
       US 6150508
                                19980318 (9)
ΑI
       US 1998-44668
       Continuation-in-part of Ser. No. US 1997-827017, filed on 25 Mar 1997,
RLI
       now abandoned which is a continuation-in-part of Ser. No. US
       1996-621399, filed on 25 Mar 1996, now abandoned
DT
       Utility
FS
       Granted
       Primary Examiner: Ungar, Susan
EXNAM
       Townsend and Townsend and Crew LLP
LREP
       Number of Claims: 16
CLMN
       Exemplary Claim: 1,2,7,12
ECL
DRWN
       24 Drawing Figure(s); 20 Drawing Page(s)
LN.CNT 1896
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention relates to monoclonal antibodies that bind to the
AB
       extracellular domain of prostate-specific membrane antigen (PSMA),
       hybridoma cell lines producing the antibodies, and methods of using such
       antibodies for diagnosis and treatment of cancer. In particular,
       thirty-five monoclonal antibodies reactive with PSMA expressed on the
       cell surface are exemplified. Additionally, the present invention
       relates to a novel protein variant (PSM') of PSMA detected by a number
       of the antibodies of the invention. The hydrolase activity of PSMA and
       PSM' allows the use of an immunoenzymatic assay for their detection.
     ANSWER 9 OF 24 USPATFULL
L6
       2000:70963 USPATFULL
AN
       Reshaped monoclonal antibodies against an immunoglobulin
TI
       isotype
       Hardman, Norman, Riehen, Switzerland
IN
       Kolbinger, Frank, Freiburg, Germany, Federal Republic of
       Saldanha, Jose, Enfield, United Kingdom
PA
       Novartis Corporation, Summit, NJ, United States (U.S. corporation)
```

Tanox Biosystems, Inc., Houston, TX, United States (U.S. corporation)

```
PΤ
       US 6072035
                                 20000606
       US 1995-485246
                                 19950607 (8)
ΑI
       Division of Ser. No. US 1993-127721, filed on 27 Sep 1993 which is a
RLI
       continuation-in-part of Ser. No. US 1992-952802, filed on 25 Sep 1992,
       now abandoned
DT
       Utility
FS
       Granted
       Primary Examiner: Reeves, Julie
EXNAM
       Ferraro, Gregory D.
LREP
       Number of Claims: 20
CLMN
       Exemplary Claim: 1
ECL
       2 Drawing Figure(s); 1 Drawing Page(s)
DRWN
LN.CNT 2732
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention relates to reshaped human monoclonal antibodies directed
ΑB
       against isotypic determinants of immunoglobulin E (IgE),
       direct equivalents and derivatives of said antibodies. The molecules of
       the invention are useful for diagnostics, prophylaxis and treatment of
       allergy.
     ANSWER 10 OF 24 USPATFULL
L6
ΑN
       2000:64943 USPATFULL
       Reshaped monoclonal antibodies against an immunoglobulin
ΤI
       isotype
       Hardman, Norman, Riehen, Switzerland
ΙN
       Kolbinger, Frank, Freiburg, Germany, Federal Republic of
       Saldanha, Jose, Enfield, United Kingdom
       Novartis Corporation, Summit, NJ, United States (U.S. corporation)
Tanox Biosystems, Inc., Houston, TX, United States (U.S. corporation)
PA
                                  20000523
PΙ
       US 6066718
ΑI
       US 1993-127721
                                  19930927 (8)
       Continuation-in-part of Ser. No. US 1992-952802, filed on 25 Sep 1992,
RLI
       now abandoned
DT
       Utility
FS
       Granted
EXNAM Primary Examiner: Hutzell, Paula K.; Assistant Examiner: Worrall,
       Timothy A.
LREP
       Ferrar, Gregory D.
       Number of Claims: 10
CLMN
ECL
       Exemplary Claim: 1
       1 Drawing Figure(s); 1 Drawing Page(s)
DRWN
LN.CNT 2830
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention relates to reshaped human monoclonal antibodies directed
AR
       against isotypic determinants of immunoglobulin E (IgE),
       direct equivalents and derivatives of said antibodies. The molecules of
       the invention are useful for diagnostics, prophylaxis and treatment of
       allergy.
L6
     ANSWER 11 OF 24 USPATFULL
AN
        2000:27563 USPATFULL
       Method for detecting the presence of P-selectin
TT
       Chesnut, Robert W., Cardiff, CA, United States
TN
        Polley, Margaret J., La Jolla, CA, United States
        Paulson, James C., Del Mar, CA, United States
        Jones, S. Tarran, Radlett, United Kingdom
        Saldanha, Jose W., Middlesex, United Kingdom
        Bendig, Mary M., London, United Kingdom
        Kriegler, Michael, Rancho Santa Fe, CA, United States
        Perez, Carl, San Diego, CA, United States
       Bayer, Robert, San Diego, CA, United States
Nunn, Michael, San Diego, CA, United States
Cytel Corporation, San Diego, CA, United States (U.S. corporation)
PA
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20000307

PΙ

US 6033667

19971105 (8) US 1997-964690 AΙ Continuation of Ser. No. US 1994-202047, filed on 25 Feb 1994, now RLI patented, Pat. No. US 5800815 which is a continuation-in-part of Ser. No. US 1993-57292, filed on 5 May 1993, now abandoned which is a continuation-in-part of Ser. No. US 1992-880196, filed on 5 May 1992, now abandoned IL 1993-105614 19930505 PRAI DTUtility FS Granted **EXNAM** Primary Examiner: Chan, Christina Y.; Assistant Examiner: Gambel, Phillip LREP Campbell & Flores LLP CLMN Number of Claims: 21 ECL Exemplary Claim: 1 DRWN 44 Drawing Figure(s); 40 Drawing Page(s) LN.CNT 4009 CAS INDEXING IS AVAILABLE FOR THIS PATENT. The present invention relates to compositions and methods for treating AΒ inflammation and other pathological conditions using novel blocking P-selectin antibodies that inhibit adhesion of leukocytes to activated platelets and/or to activated vascular endothelium in vivo. Both murine and humanized antibodies are provided. L6 ANSWER 12 OF 24 USPATFULL 2000:12616 USPATFULL ΑN TΙ Chimeric antibodies Hardman, Norman, Riehen, Switzerland ΤN Gill, Laura Lee, Riehen, Switzerland de Winter, Ronald F. J., Milton Ernest, United Kingdom Wagner, Kathrin, Basel, Switzerland Heusser, Christoph, Bottmingen, Switzerland Ciba-Geigy Corporation, Tarrytown, NY, United States (U.S. corporation) PA PΙ US 6020153 20000201 ΑI US 1994-307087 19940916 (8) Continuation of Ser. No. US 1992-947897, filed on 18 Sep 1992, now RLI abandoned which is a continuation of Ser. No. US 1988-287178, filed on 21 Dec 1988, now abandoned PRAI GB 1988-77 19880105 19880824 GB 1988-20099 DTUtility FS Granted Primary Examiner: Ziska, Suzanne E. EXNAM Nowak, Henry P., Elmer, James Scott, Foley, Shawn P. LREP CLMN Number of Claims: 25 ECL Exemplary Claim: 1 DRWN 7 Drawing Figure(s); 7 Drawing Page(s) LN.CNT 2592 CAS INDEXING IS AVAILABLE FOR THIS PATENT. AB The invention relates to murine/human chimeric monoclonal antibodies with high specificity to and affinity for human carcinoembryonic antigen (CEA), derivatives thereof, processes for the ${\bf preparation}$ of these antibodies and their derivatives, DNAs coding for heavy and light chains of these antibodies, processes for the preparation of said DNAs, mammalian cell lines that produce and secrete the antibodies and processes for the preparation of said cell lines. The chimeric antibodies and their derivatives are used for clinical purposes in vitro and in vivo, especially for the diagnosis of cancer, for localization and in vivo imaging of tumors, for therapy, e.g. site-directed delivery of cytotoxins, and similar purposes. The invention also concerns test kits and pharmaceutical compositions containing said chimeric monoclonal antibodies and/or derivatives thereof.

L6

```
2000:10021 USPATFULL
ΑN
       Antibody against human interleukin-5-receptor .alpha. chain
TI
       Koike, Masamichi, Tokyo, Japan
TN
       Furuya, Akiko, Tokyo, Japan
       Nakamura, Kazuyasu, Tokyo, Japan
       Iida, Akihiro, Tokyo, Japan
       Anazawa, Hideharu, Tokyo, Japan
       Hanai, Nobuo, Kanagawa, Japan
       Takatsu, Kiyoshi, Tokyo, Japan
       Kyowa Hakko Kogyo Co., Ltd., Tokyo, Japan (non-U.S. corporation)
PA
       US 6018032
                                20000125
PI
       WO 9710354 19970320
       US 1997-836561
                                19970509 (8)
AΙ
       WO 1996-JP2588
                                19960911
                                19970509 PCT 371 date
                                19970509 PCT 102(e) date
       JP 1995-232384
                            19950911
PRAI
DT
       Utility
FS
       Granted
       Primary Examiner: Mertz, Prema
EXNAM
       Pennie & Edmonds LLP
LREP
CLMN
       Number of Claims: 13
       Exemplary Claim: 1
ECL
DRWN
       61 Drawing Figure(s); 61 Drawing Page(s)
LN.CNT 5703
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention provides monoclonal antibodies and humanized
       antibodies which react specifically with a human interleukin-5 receptor
       .alpha. chain. The invention also provides hybridomas and transformants
       which produce the antibodies, the monoclonal antibodies and humanized
       antibodies, a method for detecting an interleukin-5 receptor .alpha.
       chain immunologically by means of these antibodies, as well as a method
       for diagnosing and treating diseases such as chronic bronchial asthma by
       means of the monoclonal antibodies and humanized antibodies. The present
       invention is useful for diagnosis or treatment of diseases such as
       chronic bronchial asthma.
     ANSWER 14 OF 24 USPATFULL
L6
       1999:136684 USPATFULL
AN
       Inhibition of intimal hyperplasia using antibodies to PDGF receptors and
TΙ
       heparin
       Hart, Charles E., Brier, WA, United States
Kenagy, Richard D., Seattle, WA, United States
IN
       Clowes, Alexander W., Seattle, WA, United States
       ZymoGenetics, Inc., Seattle, WA, United States (U.S. corporation)
PΑ
       US 5976534
                                19991102
PΙ
       US 1995-482533 19950607 (8)
Continuation-in-part of Ser. No. US 1994-366860, filed on 30 Dec 1994,
ΑI
RLI
       now patented, Pat. No. US 5620687 which is a continuation-in-part of
       Ser. No. US 1994-304623, filed on 12 Sep 1994, now abandoned which is a
       continuation of Ser. No. US 1993-23504, filed on 25 Feb 1993, now
       abandoned
DT
       Utility
FS
       Granted
       Primary Examiner: Feisee, Lila; Assistant Examiner: Gambel, Phillip
EXNAM
       Parker, Gary E.
LREP
       Number of Claims: 37
CLMN
ECL
       Exemplary Claim: 1
       12 Drawing Figure(s); 12 Drawing Page(s)
DRWN
LN.CNT 2864
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Methods for inhibiting intimal hyperplasia in the vasculature of
AB
       mammals, including primates, are disclosed. The methods comprise
```

administering to the mammal an anti-PDGF receptor antibody, such as an

anti-PDGF-alpha receptor antibody or an anti-PDGF-beta receptor antibody. The methods are useful in reducing intimal hyperplasia due to, for example, vascular injuries resulting from angioplasty, endarterectomy, reduction atherectomy or anastomosis of a vascular graft. The anti-PDGF receptor antibodies may optionally be administered coordinately with heparin, whereby the coordinately administered antibody and heparin are combinatorially effective in inhibiting intimal hyperplasia.

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L6
     ANSWER 15 OF 24 USPATFULL
       1999:117280 USPATFULL
ΑN
       Reshaped monoclonal antibodies against an immunoglobulin
TI
       isotype
IN
       Hardman, Norman, Riehen, Switzerland
       Kolbinger, Frank, Freiburg, Germany, Federal Republic of
       Saldanha, Jose, Enfield, United Kingdom
       Novartis Corporation, Summit, NJ, United States (U.S. corporation)
Tanox Biosystems, Inc., Houston, TX, United States (U.S. corporation)
PΑ
PΙ
       US 5958708
                                19990928
       US 1995-476176
ΑI
                                19950607 (8)
       Division of Ser. No. US 1993-127721, filed on 27 Sep 1993 which is a
RLI
       continuation-in-part of Ser. No. US 1992-952802, filed on 25 Sep 1992,
       now abandoned
DT
       Utility
FS
       Granted
       Primary Examiner: Caputa, Anthony C.; Assistant Examiner: Navarro, Mark
EXNAM
       Ferraro, Gregory D.
LREP
       Number of Claims: 8
CLMN
       Exemplary Claim: 1
ECL
       1 Drawing Figure(s); 1 Drawing Page(s)
DRWN
LN.CNT 2666
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention relates to reshaped human monoclonal antibodies directed
AΒ
       against isotypic determinants of immunoglobulin E (IgE),
       direct equivalents and derivatives of said antibodies. The molecules of
       the invention are useful for diagnostics, prophylaxis and treatment of
       allergy.
     ANSWER 16 OF 24 USPATFULL
L6
       1998:159916 USPATFULL
AN
       Method of enhancing proliferation or differentiation of hematopoietic
TΙ
       stem cells using Wnt polypeptides
       Matthews, William, Woodside, CA, United States
ΙN
       Austin, Timothy W., Morgan Hill, CA, United States
       Genentech, Inc., South San Francisco, CA, United States (U.S.
PΑ
       corporation)
PΙ
       US 5851984
                                 19981222
ΑI
       US 1996-696566
                                19960816 (8)
DT
       Utility
FS
       Granted
       Primary Examiner: Fitzgerald, David L.; Assistant Examiner: Basham,
EXNAM
       Darvl A.
       Svoboda, Craig G., Marschang, Diane L.
LREP
CLMN
       Number of Claims: 20
ECL
       Exemplary Claim: 1
DRWN
       4 Drawing Figure(s); 2 Drawing Page(s)
LN.CNT 3923
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Uses for Wnt polypeptides in hematopoiesis are disclosed. In particular,
AB
       in vitro and in vivo methods for enhancing proliferation or
       differentiation of a hematopoietic stem/progenitor cell using a Wnt
       polypeptide, and optionally another cytokine, are described.
```

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1998:153858 USPATFULL
AN
       Antibodies to the antigen campath-1
ΤI
       Waldmann, Herman, Cambridge, United Kingdom
IN
       Clark, Michael R., Cambridge, United Kingdom
       Winter, Gregory P., Cambridge, United Kingdom
       Riechmann, Lutz, La Jolla, CA, United States
       British Technology Group Limited, London, United Kingdom (non-U.S.
PA
       corporation)
PΙ
       US 5846534
                               19981208
ΑI
       US 1994-235705
                               19940429 (8)
       Continuation of Ser. No. US 1993-99480, filed on 30 Jul 1993, now
RLI
       abandoned which is a continuation of Ser. No. US 1992-921601, filed on 3
       Aug 1992, now abandoned which is a continuation of Ser. No. US
       1989-424233, filed on 12 Oct 1989, now abandoned
PRAI
       GB 1988-3228
                           19880212
       GB 1988-4464
                           19880225
DT
       Utility
FS
       Granted
       Primary Examiner: Budens, Robert D.
EXNAM
LREP
       Nixon & Vanderhye
       Number of Claims: 18
CLMN
       Exemplary Claim: 1,13
ECL
DRWN
       21 Drawing Figure(s); 13 Drawing Page(s)
LN.CNT 1094
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
AB
       An antibody is produced, which will bind effectively with the antigen
       Campath-1, and which has at least one complementarity determining region
       of rat origin, as identified in FIG. 2, which may be combined with a
       range of different foreign variable domain framework regions as desired,
       including framework regions of human origin.
     ANSWER 18 OF 24 USPATFULL
L6
       1998:150724 USPATFULL
ΑN
TΙ
       Chimeric antibodies
IN
       Hardman, Norman, Riehen, Switzerland
       Gill, Laura Lee, Riehen, Switzerland
       de Winter, Ronald F.J., Milton Ernest, England
       Wagner, Kathrin, Basel, Switzerland
       Heusser, Christoph, Bottmingen, Switzerland
       CIBA-GEIGY Corporation, Tarrytown, NY, United States (U.S. corporation)
PA
PΙ
       US 5843708
                                19981201
       US 1995-462371
                                19950605 (8)
ΑT
       Division of Ser. No. US 1994-307087, filed on 16 Sep 1994 which is a
RLI
       continuation of Ser. No. US 1992-947897, filed on 18 Sep 1992, now
       abandoned which is a continuation of Ser. No. US 1988-287178, filed on
       21 Dec 1988, now abandoned
       GB 1988-77
                           19880105
PRAI
       GB 1988-20099
                           19880824
חת
       Utility
FS
       Granted
EXNAM
       Primary Examiner: Ziska, Suzanne E.
LREP
       Nowak, Henry P.
       Number of Claims: 21
CLMN
ECL
       Exemplary Claim: 1
DRWN
       9 Drawing Figure(s); 7 Drawing Page(s)
LN.CNT 2247
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention relates to murine/human chimeric monoclonal antibodies
AB
       with high specificity to and affinity for human carcinoembryonic antigen
       (CEA), derivatives thereof, processes for the preparation of
       these antibodies and their derivatives, DNAs coding for heavy and light
       chains of these antibodies, processes for the preparation of
       said DNAs, mammalian cell lines that produce and secrete the antibodies
```

and processes for the preparation of said cell lines. The

chimeric antibodies and their derivatives are used for clinical purposes in vitro and in vivo, especially for the diagnosis of cancer, for localization and in vivo imaging of tumors, for therapy, e.g. site-directed delivery of cytotoxins, and similar purposes. The invention also concerns test kits and pharmaceutical compositions containing said chimeric monoclonal antibodies and/or derivatives thereof.

```
L6
     ANSWER 19 OF 24 USPATFULL
ΑN
       1998:104391 USPATFULL
       Antibodies to P-selectin and their uses
TΙ
       Chestnut, Robert W., Cardiff, CA, United States Polley, Margaret J., La Jolla, CA, United States
ΙN
       Paulson, James C., Del Mar, CA, United States
       Jones, S. Tarran, Radlett, United Kingdom
       Saldanha, Jose W., Middlesex, United Kingdom
       Bendig, Mary M., London, United Kingdom
       Kriegler, Michael, Rancho Santa Fe, CA, United States
       Perez, Carl, San Diego, CA, United States
       Bayer, Robert, San Diego, CA, United States
Nunn, Michael, San Diego, CA, United States
       Cytel Corporation, San Diego, CA, United States (U.S. corporation)
PA
PΙ
       US 5800815
                                 19980901
       US 1994-202047 19940225 (8)
Continuation-in-part of Ser. No. US 1993-57292, filed on 5 May 1993, now
ΑI
RLI
       abandoned which is a continuation-in-part of Ser. No. US 1992-880198,
       filed on 5 May 1992, now abandoned
                            19030505
       IL 1903-105614
PRAI
                            19930504
       WO 1993-US4274
       Utility
DТ
FS
       Granted
       Primary Examiner: Hutzell, Paula K.; Assistant Examiner: Gambel, Phillip
EXNAM
       Campbell & Flores LLP
LREP
       Number of Claims: 57
CLMN
       Exemplary Claim: 32,40
ECL
       47 Drawing Figure(s); 40 Drawing Page(s)
DRWN
LN.CNT 4013
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention relates to compositions and methods for treating
AΒ
       inflammation and other pathological conditions using novel blocking
       P-selectin antibodies that inhibit adhesion of leukocytes to activated
       platelets and/or to activated vascular endothelium in vivo. Both murine
       and humanized antibodies are provided.
     ANSWER 20 OF 24 USPATFULL
L6
       1998:82874 USPATFULL
ΑN
       Monoclonal antibodies to cytotoxic lymphocyte maturation factor
TΙ
       Gately, Maurice Kent, Montville, NJ, United States
IN
       Gubler, Ulrich Andreas, Glen Ridge, NJ, United States
       Hulmes, Jeffrey David, Ringwood, NJ, United States
       Podlaski, Frank John, New City, NY, United States
       Stern, Alvin Seth, Passaic Park, NJ, United States
       Chizzonite, Richard Anthony, South Kent, CT, United States
       Pan, Yu-Ching Eugene, Pine Brook, NJ, United States
       Hoffmann-La Roche Inc., Nutley, NJ, United States (U.S. corporation)
PA
                                 19980714
PΤ
       US 5780597
                                 19950602 (8)
       US 1995-460061
ΑI
       Division of Ser. No. US 1994-205011, filed on 2 Mar 1994, now abandoned
RLI
       which is a division of Ser. No. US 1992-857023, filed on 24 Mar 1992,
       now abandoned which is a continuation-in-part of Ser. No. US
       1990-572284, filed on 27 Aug 1990, now abandoned which is a
       continuation-in-part of Ser. No. US 1990-520935, filed on 9 May 1990,
       now abandoned which is a continuation-in-part of Ser. No. US
       1989-455708, filed on 22 Dec 1989, now abandoned
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DΤ
       Utility
FS
       Granted
       Primary Examiner: Cunningham, Thomas M.; Assistant Examiner: Lubet,
EXNAM
       Johnston, George W., Epstein, William H., Buchholz, Briana C.
LREP
       Number of Claims: 3
CLMN
       Exemplary Claim: 1
ECL
DRWN
       41 Drawing Figure(s); 44 Drawing Page(s)
LN.CNT 2912
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention relates to antibodies which bind to a novel
AΒ
       cytotoxic lymphocyte maturation factor. When bound to the cytotoxic
       lymphocyte maturation factor, the antibodies can neutralize bioactivity
       of the factor.
     ANSWER 21 OF 24 USPATFULL
L6
       97:118165 USPATFULL
ΑN
       Product and process for targeting an immune response
TΙ
       Nemazee, David A., Denver, CO, United States
IN
       National Jewish Center for Immunology and Respiratory Medicine, Denver,
PA
       CO, United States (U.S. corporation)
       US 5698679
                                19971216
PΙ
ΑI
       US 1994-309006
                                19940919 (8)
DT
       Utility
FS
       Granted
       Primary Examiner: Feisee, Lila; Assistant Examiner: Eyler, Yvonne
EXNAM
LREP
       Sheridan Ross P.C.
CLMN
       Number of Claims: 27
       Exemplary Claim: 1
ECL
       5 Drawing Figure(s); 5 Drawing Page(s)
DRWN
LN.CNT 1793
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention relates to a product and process for regulating an
AB
       immune system using an immunoglobulin fusion protein capable
       of targeting a specific peptide precursor to a specific antigen
       presenting cell. Disclosed is a peptide precursor associated with an
       immunoglobulin molecule capable of binding to an antigen on the
       surface of an antigen presenting cell. Also disclosed is a nucleic acid
       molecule having a sequence encoding an immunoglobulin fusion
       protein comprising a peptide precursor and an immunoglobulin
       molecule. The invention is additionally directed to therapeutic reagents
       which can act as toleragens or immunogens useful in the regulation of an
       immune response.
     ANSWER 22 OF 24 USPATFULL
L6
       97:31409 USPATFULL
ΑN
       Inhibition of intimal hyperplasia using antibodies to PDGF beta
ΤI
       receptors
       Hart, Charles E., Brier, WA, United States
TN
       Kenagy, Richard D., Seattle, WA, United States
       Clowes, Alexander W., Seattle, WA, United States
       ZymoGenetics, Inc., Seattle, WA, United States (U.S. corporation)
PA
       University of Washington, Seattle, WA, United States (U.S. corporation)
                                19970415
PΙ
       US 5620687
       US 1994-366860 19941230 (8)
Continuation-in-part of Ser. No. US 1994-304623, filed on 12 Sep 1994,
ΑI
RLI
       now abandoned which is a continuation of Ser. No. US 1993-23504, filed
       on 25 Feb 1993, now abandoned
DT
       Utility
FS
       Granted
       Primary Examiner: Chan, Christina Y.; Assistant Examiner: Gambel,
EXNAM
       Phillip
       Parker, Gary E., Leith, Debra K., Sawislak, Deborah A.
LREP
CLMN
       Number of Claims: 19
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Exemplary Claim: 1
ECL
DRWN
       12 Drawing Figure(s); 12 Drawing Page(s)
LN.CNT 2786
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Methods for inhibiting intimal hyperplasia in the vasculature of
ΑB
       mammals, including primates, are disclosed. The methods comprise
       administering to the mammal an anti-PDGF receptor antibody, such as an
       anti-PDGF-alpha receptor antibody or an anti-PDGF-beta receptor
       antibody. The methods are useful in reducing intimal hyperplasia due to,
       for example, vascular injuries resulting from angioplasty,
       endarterectomy, reduction atherectomy or anastomosis of a vascular
       graft. The anti-PDGF receptor antibodies may optionally be administered
       coordinately with heparin, whereby the coordinately administered
       antibody and heparin are combinatorially effective in inhibiting intimal
       hyperplasia.
    ANSWER 23 OF 24 USPATFULL
L6
       97:6049 USPATFULL
ΑN
ΤI
       Method of refolding human IL-13
       Culpepper, Janice, Mountain View, CA, United States
IN
       McKenzie, Andrew, Redwood City, CA, United States
       Dang, Warren, San Jose, CA, United States
       Zurawski, Gerard, Redwood City, CA, United States
PΑ
       Schering Corporation, Kenilworth, NJ, United States (U.S. corporation)
PΙ
       US 5596072
                               19970121
ΑI
       US 1993-12543
                               19930201 (8)
       Continuation-in-part of Ser. No. US 1992-933416, filed on 21 Aug 1992,
RLI
       now abandoned
       Utility
DΤ
       Granted
FS
EXNAM
       Primary Examiner: Draper, Garnette D.; Assistant Examiner: Spector,
       Lorraine M.
       Ching, Edwin P.
LREP
       Number of Claims: 10
CLMN
ECL
       Exemplary Claim: 1
DRWN
       288 Drawing Figure(s); 61 Drawing Page(s)
LN.CNT 4619
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Nucleic acids encoding human IL-13, and purified IL-13 proteins and
AΒ
       fragments thereof. Antibodies, both polyclonal and monoclonal, are also
       provided. Methods of using the compositions for both diagnostic and
       therapeutic utilities are provided.
     ANSWER 24 OF 24 USPATFULL
L6
       97:3527 USPATFULL
AN
ΤI
       Method of producing an anti-D immunoglobulin
       concentrate and a pharmaceutical preparation
ΙN
       Hodler, Gerhard, Worb, Switzerland
       Lerch, Peter, Bern, Switzerland
       Stucki, Martin, Laupen, Switzerland
       Rotkreuzstiftung Zentrallaboratorium Blutspendedienst, Bern, Switzerland
PΑ
       (non-U.S. corporation)
       US 5593675
                                19970114
PΤ
AΤ
       US 1994-360334
                                19941221 (8)
PRAI
       CH 1993-93810912
                           19931227
DT
       Utility
FS
       Granted
EXNAM
       Primary Examiner: Feisee, Lila
LREP
       Seed and Berry LLP
CLMN
       Number of Claims: 20
ECL
       Exemplary Claim: 1
DRWN
       1 Drawing Figure(s); 1 Drawing Page(s)
LN.CNT 552
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AΒ An anti-D immunoglobulin G-preparation is produced in high-yield from human plasma containing anti-D IgG, or a plasma fraction containing an anti-D IgG, in which (A) the plasma, or the plasma fraction, with a pH in the range of pH 3.5 to 6.5and a conductivity value in the range of 2 to 4 mS/cm, is subjected to ion exchange chromatography with an adsorbent which has carboxymethyl groups as functional groups, the anti-D IgG being bound to the adsorbent, (B) the adsorbent with the bound anti-D IgG is first rinsed with a wash solution at a pH in the range of 5 to 8 and a conductivity value in the range of 2 to 4 mS/cm, and the anti-D IgG is subsequently eluted, and further (C) the eluted anti-D IgG at a pH in the range of 6 to 8 and a conductivity value in the range of 2 to 4 mS/cm is treated with an alkaline adsorbent with ion-exchange characteristics in order to bind undesired components, and finally, the anti-D IgG is concentrated. The non-infectious anti-D concentrate thus obtained possesses a high specific activity of more than 1% anti-D IgG per gram of total IgG.

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s concentr? and lyophiliz? and immunoglobulin?
       196725 CONCENTR?
        60084 CONC
        24730 CONCS
        75545 CONC
                 (CONC OR CONCS)
       133608 CONCD
         12528 CONCG
       1273077 CONCN
       815460 CONCNS
       1794486 CONCN
                 (CONCN OR CONCNS)
       2003854 CONCENTR?
                 (CONCENTR? OR CONC OR CONCD OR CONCG OR CONCN)
        12971 LYOPHILIZ?
        70372 IMMUNOGLOBULIN?
         40048 IG
        11178 IGS
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                 (IG OR IGS)
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                 (IMMUNOGLOBULIN? OR IG)
1.7
            77 CONCENTR? AND LYOPHILIZ? AND IMMUNOGLOBULIN?
=> d ibib all
    ANSWER 1 OF 77 HCAPLUS COPYRIGHT 2001 ACS
                         2000:683899 HCAPLUS
ACCESSION NUMBER:
                         134:67869
DOCUMENT NUMBER:
                         Fibrinolytic serine protease from Spirodela polyrhiza
TITLE:
                         Choi, Hye-Seon
AUTHOR(S):
                         Department of Biological Sciences, University of
CORPORATE SOURCE:
                         Ulsan, Ulsan, 680-749, S. Korea
                         KORUS'99, Proc. Russ.-Korean Int. Symp. Sci. Technol.,
SOURCE:
                         3rd (1999), Volume 2, 437-440. Novosibirsk State
                         Technical University: Novosibirsk, Russia.
                         CODEN: 69AKRG
DOCUMENT TYPE:
                         Conference
                         English
LANGUAGE:
    2000:683899 HCAPLUS
AN
DN
    134:67869
TΙ
    Fibrinolytic serine protease from Spirodela polyrhiza
    Choi, Hye-Seon
ΑIJ
    Department of Biological Sciences, University of Ulsan, Ulsan, 680-749, S.
CS
    Korea
    KORUS'99, Proc. Russ.-Korean Int. Symp. Sci. Technol., 3rd (1999), Volume
SO
    2, 437-440 Publisher: Novosibirsk State Technical University, Novosibirsk,
    Russia.
    CODEN: 69AKRG
DΤ
    Conference
LA
    English
CC
    7-2 (Enzymes)
    A serine protease was purified from Chinese herb (Spirodela polyrhiza).
AΒ
    Protease has a mol. mass of 180,000 dalton and 43,000 dalton in gel
     filtration and SDS-PAGE, resp., implying it is a trimer. Its optimum pH
    was 4.5-5.0. Enzyme was stable below 40C.degree. and after
    lyophilization. The enzyme activity was inhibited significantly
    by leupeptin and to a lesser degree by PMSF. The protease hydrolyzed not
    only fibrin but also fibrinogen, cleaving A and B without affecting chain
    of fibrinogen. However, no hydrolysis was found with Hb, Ig,
    and albumin under the same condition. It cleaved preferentially Arg or
    Lys residue of synthetic substrates (P' position). The enzyme had an
    anticoagulatory activity measured with activated partial thromboplastin
    time and thrombin time test. It delayed APTT and TT two times at the
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protein concn. of 5.0 and 5.7 ug, resp. and drastically reduced
     after heat treatment.
     serine proteinase fibrinolysis Spirodela
ST
     Anticoagulants
TT
        (fibrinolytic serine protease from Spirodela polyrhiza)
TΤ
     Fibrinogens
     Fibrins
     RL: BPR (Biological process); BIOL (Biological study); PROC (Process)
        (fibrinolytic serine protease from Spirodela polyrhiza)
IT
     37259-58-8P, Serine proteinase
     RL: BAC (Biological activity or effector, except adverse); BPR (Biological
     process); PRP (Properties); PUR (Purification or recovery); BIOL
     (Biological study); PREP (Preparation); PROC (Process)
        (fibrinolytic serine protease from Spirodela polyrhiza)
                            65316-83-8
                                          69861-90-1
                                                        73392-19-5
     9002-04-4, Thrombin
TΤ
     RL: BPR (Biological process); BIOL (Biological study); PROC (Process)
        (fibrinolytic serine protease from Spirodela polyrhiza)
RE.CNT
RE
(1) Choi, H; Mycologia 1998, V90, P674 HCAPLUS
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(5) Nikai, T; Arch Biochem Biophys 1984, V231, P309 HCAPLUS
(6) Noeske-Jungblut, C; J Biol Chem 1995, V270, P28629 HCAPLUS
(7) Olson, S; J Biol Chem 1992, V267, P12528 HCAPLUS
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=> d 2-77
     ANSWER 2 OF 77 HCAPLUS COPYRIGHT 2001 ACS
1.7
     2000:592533 HCAPLUS
AN
     133:198650
DN
     Biodegradable composite material for the production of microcapsules
TΙ
     Teller, Marianne; Heinrich, Hans-Werner; Teller, Joachim; Meyer, Udo
ΙN
     Bioserv A.-G., Germany
PΑ
     PCT Int. Appl., 24 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
     German
FAN.CNT 1
                            DATE
                                             APPLICATION NO.
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     WO 2000048573
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             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                           DE 2000-10008880 20000220
                             20000824
     DE 10008880
                        A1
PRAI DE 1999-19907227 19990219
RE.CNT 6
RE
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(5) Takeda Chemical Industries Ltd; EP 0601799 A 1994 HCAPLUS
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L7
ΑN
     2000:149842 HCAPLUS
DN
    132:150931
    Manufacture of milk power with specific immune globulin
ΤI
ΙN
    Wang, Wenrong
PA
     Peop. Rep. China
     Faming Zhuanli Shenging Gongkai Shuomingshu, 5 pp.
SO
     CODEN: CNXXEV
DT
    Patent
LA
    Chinese
FAN.CNT 1
     PATENT NO. KIND DATE
                                       APPLICATION NO. DATE
                                      -----
    CN 1173288 A 19980218
                                       CN 1997-115852 19970828
PΙ
                    В 20001206
    CN 1059128
L7
    ANSWER 4 OF 77 HCAPLUS COPYRIGHT 2001 ACS
    2000:65297 HCAPLUS
AN
DN
    132:113093
    Compositions containing lysin enzyme for prevention and treatment of group
ΤI
    A streptococcal infections
IN
    Fischetti, Vincent; Loomis, Lawrence
    New Horizons Diagnostics, USA
PA
SO
    U.S., 6 pp., Cont.-in-part of U.S. Ser. No. 962,523.
    CODEN: USXXAM
DT
    Patent
LA
    English
FAN.CNT 3
                                 APPLICATION NO. DATE
    PATENT NO. KIND DATE
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    US 6017528 A 20000125
                                       US 1999-257025 19990225
PΙ
    US 5997862
                    Α
                         19991207
                                       US 1997-962523 19971031
                                   WO 1999-US4063 19990225
    WO 2000050069 A1 20000831
            AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
            DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP,
            KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN,
            MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
            TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU,
            TJ, TM
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                                      AU 1999-28767 19990225
    AU 9928767
                    A1
                          20000914
PRAI US 1997-962523
                    19971031
    US 1999-257025
                    19990225
    WO 1999-US4063 19990225
    ANSWER 5 OF 77 HCAPLUS COPYRIGHT 2001 ACS
L7
AN 
    1999:732954 HCAPLUS
DN
    131:356096
    Oral delivery system containing a group C streptococcal phage-associated
TΙ
    lysin enzyme for prophylactic and therapeutic treatment of group A
    streptococcal infection
ΙN
    Fischetti, Vincent; Loomis, Lawrence
PΑ
    New Horizons Diagnostics Corp., USA
SO
    U.S., 5 pp., Division of U.S. Ser. No. 962,523.
    CODEN: USXXAM
DT
    Patent
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    English
FAN.CNT 3
                KIND DATE
    PATENT NO.
                                       APPLICATION NO. DATE
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US 1999-257026
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PΙ
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PRAI US 1997-962523
                      19971031
     WO 1999-US4063
                      19990225
RE.CNT 2
RE
(1) Fischetti; US 5604109 1997 HCAPLUS
(2) Scott; US 4784948 1988 HCAPLUS
L7
     ANSWER 6 OF 77 HCAPLUS COPYRIGHT 2001 ACS
ΑN
     1999:182963 HCAPLUS
     130:219694
DN
     A new enzyme immunoassay for soluble fibrin in plasma, with a high
ΤI
     discriminating power for thrombotic disorders
     Bos, R.; Laterveer-Vreeswijk, G. H.; Lockwood, D.; Szewczyk, K.;
ΑU
     Nieuwenhuizen, W.
     Gaubius Laboratory, TNO Prevention Health, Leiden, 2333 CA, Neth.
CS
     Thromb. Haemostasis (1999), 81(1), 54-59
SO
     CODEN: THHADQ; ISSN: 0340-6245
     F. K. Schattauer Verlagsgesellschaft mbH
PΒ
DT
     Journal
LA
     English
RE.CNT 30
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(6) Dempfle, C; Thromb Haemost 1995, V74, P673 HCAPLUS
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     ANSWER 7 OF 77 HCAPLUS COPYRIGHT 2001 ACS
L7
     1999:172617 HCAPLUS
ΑN
     130:213644
DN
ΤI
     Dried biologically or therapeutically active preparations
     Kanellos, Jerry; Oates, Adrian; Goss, Neil
ΙN
PA
     CSL Limited, Australia
SO
     PCT Int. Appl., 28 pp.
     CODEN: PIXXD2
DT
     Patent
LA
    English
FAN.CNT 1
                                           APPLICATION NO.
                      KIND
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                            _____
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                            19990304
                                          WO 1998-AU682
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     WO 9910011
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PΙ
            AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
             DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG,
             KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
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                                           ZA 1998-7633
                                                            19980824
     ZA 9807633
                            19990225
                      Α
                                           AU 1998-87231
                                                            19980825
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     AU 9887231
                       A1
                                           EP 1998-938550
                                                            19980825
                            20000621
                      A1
     EP 1009438
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
PRAI AU 1997-8719
                      19970825
                      19980825
     WO 1998-AU682
RE.CNT 10
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- (5) Miles Laboratories Inc; US 4623717 A 1986 HCAPLUS
- ALL CITATIONS AVAILABLE IN THE RE FORMAT
- L7 ANSWER 8 OF 77 HCAPLUS COPYRIGHT 2001 ACS
- AN 1998:265356 HCAPLUS
- TI Comparison of freezing and lyophilizing for preservation of colostrum as a source of immunoglobulins for calves
- AU Klobasa, F.; Goel, M. C.; Werhahn, E.
- CS Institute of Animal Husbandry and Animal Ethology, Mariensee, Germany
- SO J. Anim. Sci. (1998), 76(4), 923-926 CODEN: JANSAG; ISSN: 0021-8812
- PB American Society of Animal Science
- DT Journal
- LA English
- L7 ANSWER 9 OF 77 HCAPLUS COPYRIGHT 2001 ACS
- AN 1998:145737 HCAPLUS
- DN 128:269347
- TI Protein G affinity chromatography as a method for isolating immunoglobulins for the treatment of immunodeficiency in marine mammals
- AU Schwertner, Harvey A.; Dalton, Leslie M.; Mcbain, James F.; Patterson, Wayne R.
- CS Clinical Investigation Directorate, Wilford Hall Medical Center/RDL, Lackland AFB, TX, 78236-5319, USA
- SO Int. J. Bio-Chromatogr. (1997), 3(3), 207-214 CODEN: IJOBEQ; ISSN: 1068-0659
- PB Harwood Academic Publishers
- DT Journal
- LA English
- L7 ANSWER 10 OF 77 HCAPLUS COPYRIGHT 2001 ACS
- AN 1995:864497 HCAPLUS
- DN 123:278459
- TI Antibodies as therapeutic agents: the antivenoms
- AU Dart, Richard C.
- CS Rocky Mountain Poison Center, Denver, CO, 80204, USA
- SO J. Nat. Toxins (1995), Volume Date 1995, 4(2), 155-63 CODEN: JNTOER; ISSN: 1058-8108
- DT Journal
- LA English
- L7 ANSWER 11 OF 77 HCAPLUS COPYRIGHT 2001 ACS
- AN 1995:794883 HCAPLUS
- DN 123:179357
- TI Preparation of concentrated immunoglobulins G for therapeutic uses
- IN Burnouf, Miryana; Dernis, Dominique; Bonneel, Patrick; Burnouf, Thierry
- PA Association pour l'Essor de la Transfusion Sanguine dans la Region du Nord, Fr.
- SO Fr. Demande, 13 pp. CODEN: FRXXBL
- DT Patent
- LA French
- FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PΙ	FR 2706466	A1	19941223	FR 1993-7128	19930614
	FR 2706466	В1	19950825		
	CA 2165203	AA	19941222	CA 1994-2165203	19940613
	WO 9429334	A1	19941222	WO 1994-FR699	19940613

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W: AU, BB, BG, BR, BY, CA, CN, CZ, FI, GE, HU, JP, KE, KG, KP, KR,
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             TT, UA, US, UZ, VN
         RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE,
            BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG
                           19950103 AU 1994-70023 19940613
     AU 9470023
                      A1
                           19960403
                                          EP 1994-918916
                                                           19940613
     EP 703922
                      Α1
     EP 703922
                      В1
                           20000517
        R: AT, BE, CH, DE, DK, ES, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE
                                        JP 1994-501424 19940613
                     T2
                           19970114
     JP 09500369
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     BR 9406814
                      Α
     AT 193020
                                          AT 1994-918916
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                      Т3
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                                                           19940613
     ES 2148332
     US 6069236
                      Α
                           20000530
                                          US 1996-564030
                                                           19960325
                     19930614
PRAI FR 1993-7128
     WO 1994-FR699
                     19940613
L7
     ANSWER 12 OF 77 HCAPLUS COPYRIGHT 2001 ACS
     1995:761725 HCAPLUS
ΑN
     123:152869
DN
ΤI
     Highly concentrated immunoglobulin composition and
     method to prepare it
IN
     Eibl, Johann; Linnau, Yendra; Teschner, Wolfgang
PΑ
     Immuno A.-G., Austria
SO
     Eur. Pat. Appl., 5 pp.
     CODEN: EPXXDW
DT
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PΙ
     EP 661060
                     A2
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    Aggregation of IgG on methylated silicon surfaces studied by tapping mode
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     Preparation of immunoglobulins for intramuscular injection
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     Stepanek, Ivan
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     CODEN: CZXXA9
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    Formulation of intramuscular injections of IgG, IgM, and IgA
ΤN
    Stepanek, Ivan
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    Process for preparing an immunoglobulin preparation from
    colostrum
    Stephan, Wolfgang; Dichtelmueller, Herbert
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    Biotest Pharma G.m.b.H., Fed. Rep. Ger.
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    Eur. Pat. Appl., 5 pp.
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    Imanishi, Jiro; Ashihara, Tsukasa
    Dep. Microbiol., Kyoto Prefect. Univ. Med., Kyoto, 602, Japan
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    Cancer Res. (1990), 50(21), 7008-14
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TΙ
    Therapeutic IqM concentrates
    Collins, Michael S.; Opitz, Hans Georg; Lundblad, John L.; Seng, Richard
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    Miles, Inc., USA
SO
    Eur. Pat. Appl., 12 pp.
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     pharmaceuticals containing them
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     Von Wussow, Peter
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     Ciba-Geigy A.-G., Switz.
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     Eur. Pat. Appl., 17 pp.
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     111:239469
ΤI
    Purification and stabilization of immunoglobulin M antibodies
IN
     Dove, George; Mitra, Gautam
     Miles, Inc., USA
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$O
     Eur. Pat. Appl., 6 pp.
     CODEN: EPXXDW
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ΑN
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ΤI
    ultrāfiltration
ΑU
    Geschier, C.; Streiff, F.; Stoltz, J. F.
    Cent. Reg. Transfus. Sanquine, Vandoeuvre-les-Nancy, 51511, Fr.
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    Method for the preparation of immunoglobulins suitable for
TΤ
    intravenous administration
ΙN
    Gazzei, Guido; Giannozzi, Aldo; Valeri, Andrea
PΑ
    Sclavo S.p.A., Italy
SO
    Eur. Pat. Appl., 10 pp.
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    Method of preparing immunoglobulins against human lymphocytes
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    Hamsikova, Eva; Pardon, Jan; Ulrych, Stanislav
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PΑ
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SO
    Czech., 5 pp.
    CODEN: CZXXA9
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ΤI
    peroxide and caprylic acid
    Bulik, Jozef; Banda, Imrich; Stachy, Alfred
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SO
    Czech., 3 pp.
    CODEN: CZXXA9
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    1989:101761 HCAPLUS
DN
    110:101761
    Stable formulations of ricin toxin A chain and of its immunoconjugates and
ΤI
    stabilizer screening methods therefor
IN
    Ferris, Robert
PA
    Cetus Corp., USA
    Eur. Pat. Appl., 19 pp.
SO
    CODEN: EPXXDW
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    1988:637006 HCAPLUS
ΑN
DN
    109:237006
    Method for preparing a mixture of immunoglobulins and histamine
ΤI
    Stachy, Alfred; Bulik, Josef; Banda, Imrich
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    Czech., 2 pp.
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    1988:616033 HCAPLUS
ΑN
DN
    109:216033
    Immunoglobulin- and transferrin-containing intravenous or
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    intraperitoneal injections for control of gram-negative bacterial
    infections
    Stachy, Alfred; Lucansky, Anton; Bulik, Jozef; Banda, Imrich
IN
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    1988:498815 HCAPLUS
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    Method of producing virus- and blood group substance-free
ΤI
    immunoglobulin preparations for intravenous injection
    Uemura, Yahiro; Uriyu, Katsuhiro; Takechi, Kazuo; Hirao, Yutaka; Suyama,
ΙN
    Tadakazu
PΑ
    Green Cross Corp., Japan
    Eur. Pat. Appl., 8 pp.
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    109:79751
    Process for the preparation of lyophilized and heat-treated
TΤ
    blood-coagulation factor VIII
    Schwarz, Otto; Linnau, Yendra
IN
    Immuno A.-G. fuer Chemisch-Medizinische Produkte, Austria
PA
    Eur. Pat. Appl., 8 pp.
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    1980:220683 HCAPLUS
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DN
    92:220683
    Immunoglobulin solution suitable for intravenous use
TΤ
ΤN
    Buenning, Karl
    Blutspendedienst der Landesverbaende des Deutschen Roten Kreuzes
    Niedersachsen, Oldenburg und Bremen Gemeinnuetzige G.m.b.H., Fed. Rep.
    Ger. Offen., 12 pp.
SO
    CODEN: GWXXBX
DT
    Patent
LA
    German
FAN.CNT 1
    PATENT NO. KIND DATE
                                      APPLICATION NO. DATE
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PΙ
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                    Α
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    AT 368886
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                                         US 1980-189001
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PRAI DE 1978-2837168 19780825
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    ANSWER 65 OF 77 HCAPLUS COPYRIGHT 2001 ACS
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    1980:56611 HCAPLUS
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    92:56611
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    Stability of the lyophilized F(ab')2 fragments of horse tetanus
    antibodies isolated by affinity chromatography
    Goch, Halina; Schiller, Barbara; Korbecki, Michal
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    Serum Vaccine Res. Lab., Warsaw, 00-725, Pol.
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    Arch. Immunol. Ther. Exp. (1979), 27(4), 499-509
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    CODEN: AITEAT; ISSN: 0004-069X
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    ANSWER 66 OF 77 HCAPLUS COPYRIGHT 2001 ACS
    1979:588722 HCAPLUS
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    91:188722
    Isolation and properties of immunoperoxidase complexes
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    Gavrilova, E. M.; Dzantiev, B. B.; Egorov, A. M.
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    Dep. Chem. Enzymol., M. V. Lomonosov Moscow State Univ., Moscow, USSR
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    Biokhimiya (Moscow) (1979), 44(9), 1614-22
    CODEN: BIOHAO; ISSN: 0006-307X
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    ANSWER 67 OF 77 HCAPLUS COPYRIGHT 2001 ACS
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    1978:480234 HCAPLUS
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    89:80234
    Immunoglobulin with reduced complement binding
TΙ
    Mueller, Hans
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    Behringwerke A.-G., Ger.
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    CODEN: GWXXBX
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19771222

BR 7708559

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     FR 2375247
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                            19801128
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                       Α
                            19861128
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PRAI DE 1976-2658334 19761223
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     ANSWER 68 OF 77 HCAPLUS COPYRIGHT 2001 ACS
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     1977:451453 HCAPLUS
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     87:51453
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     An improved method for the isolation of pure IqG from human serum
TI
     Peking Metropolitan Hospital, Chronic Bronchitis Res. Team, Biochem. Dep.,
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     Peking, Peop. R. China
SO
     Sheng Wu Hua Hsueh Yu Sheng Wu Wu Li Hsueh Pao (1976), 8(3), 215-24
     CODEN: SHWPAU
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     Protein electrophoresis of liquor cerebrospinalis with cellulose acetate
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     membranes. I. Comparison of two procedures to concentrate
     proteins of collected liquor cerebrospinalis
     Kleine, T. O.; Stroh, Maria; Stroh, J.
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   . Klin.-Chem. Lab., Univ.-Nervenklin., Marburg/Lahn, Ger.
     Z. Klin. Chem. Klin. Biochem. (1974), 12(2), 66-72
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     CODEN: ZKCKAD
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     1972:445127 HCAPLUS
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     Immunochromatographic purification of human urinary luteinizing hormone
ΤI
     Van Hell, H.; Schuurs, A. H. W. M.
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     Biochem. Res. Dev. Lab., N. V. Organon, Oss, Neth.
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     J. Endocrinol. (1972), 54(1), 171-2
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     Reaction of staphylococcal protein A with rabbit immunoglobulins
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     McDowell, Graham; Grov, Arne; Oeding, Per
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     Sch. Med., Univ. Bergen, Bergen, Norway
Acta Pathol. Microbiol. Scand., Sect. B (1971), 79(6), 794-800
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     CODEN: APMIBM
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     ANSWER 72 OF 77 HCAPLUS COPYRIGHT 2001 ACS
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Preparation of freeze-dried, monomeric and immunochemically pure IgG by

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     Joustra, Marius; Lundgren, Helga
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     Res. Dep., Pharm. Fine Chem., Uppsala, Swed.
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     CODEN: PBFPA6
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     Value of autoradioimmunoelectrophoresis for studies of protein synthesis
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     by lymphocytes in vitro
     Thiele, H. G.; Stark, R.
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     I. Med. Universitaetsklin., Hamburg, Ger.
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     Z. Immunitaetsforsch., Allerg. Klin. Immunol. (1970), 140(4), 424-7
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     CODEN: ZIAIAH
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     1970:59072 HCAPLUS
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     Cleavage of human .gamma. globulin by means of cyanogen bromide
ΤI
     Sela, Michael; Arnon, Ruth; Lahav, Miriam
ΙN
     United States Dept. of Health, Education, and Welfare
PΑ
SO
     U.S., 6 pp.
     CODEN: USXXÀM
DT
     Patent
LA
     English
FAN.CNT 1
                                           APPLICATION NO.
     PATENT NO.
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                            DATE
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                                           US 1967-643165 19670602
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     US 3466368
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     ANSWER 75 OF 77 HCAPLUS COPYRIGHT 2001 ACS
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     Studies on IgA. I. (Fractionation procedure for isolation of IgA from
     pooled normal human plasma
     Zschocke, Rainer H.; Grieble, Hans G.; Bach, Gerhard L.; Anderson, Truman
ΑU
     Cook County Hosp., Chicago, Ill., USA
CS
     J. Immunol. (1969), 102(3), 625-37
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    CODEN: JOIMA3
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     Journal
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     ANSWER 76 OF 77 HCAPLUS COPYRIGHT 2001 ACS
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     1968:10857 HCAPLUS
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     Immunoelectrophoretic analysis of soluble proteins from human spleen
ΤI
     Bednarik, Tomas; Cajthamlova, H.
ΑU
     Inst. Hematol. Blood Transfusion, Prague, Czech.
CS
     (1967), 16(4), 337-41
SO
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     Journal
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     English
     ANSWER 77 OF 77 HCAPLUS COPYRIGHT 2001 ACS
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     1967:8517 HCAPLUS
ΑN
     66:8517
DN
     Recovery of long-acting thyroid stimulator from serum of patients with
TI
     thyrotoxicosis by concentration of immunoglobulin G
     Carneiro, L.; Dorrington, Keith J.; Munro, Donald S.
ΑU
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US 5102990 19920407 ΡI An IgG preparation was made by diluting 0.25 ml of Immune DETD Globulin (Human, U.S.P., Cutter Biological, which contained 15-18% protein stabilized with 0.21-0.32M glycine, with 7.25 ml of Sterile Water for Injection, U.S.P., and filtering through a 0.22 micron filter. 5 ml of the Sn (II) educing solution was mixed with 7.5 ml of the IgG preparation. The vial containing. contained the Sn (II), Sn (IV) and other salts, was discarded. The reduced and Sn (II) complexed protein fraction was concentrated by ultrafiltration to a concentration of 1.7 mg/ml. 0.5 mg aliquots of reduced and Sn (II) complexed protein were placed in sealed, N.sub.2 gas filled serum vials and frozen. A Sn (II) pertechnetate reducing solumion was made of 0.5 ml of 0.1 mM SnCl.sub.2 in 40 mM potassium biphthalat 10. . . at a pH of 5.6. The Sn (II) pertechnetate reducing solutio was added without allowing the reduced antibody solution to thaw and this solution was also frozen. A sterile, 3 mm d meter tin metal shot was added, the vial flushed with N.sub.2 and stored at -20.degree. C.. . .

L19 ANSWER 9 OF 11 USPATFULL

Polypeptides and process for the production thereof TI

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PΙ US 5089400 19920218

DETD

are injected i.p. with 2-5.times.10.sup.6 hybridoma cells. Ascitic fluid is collected repeatedly from each mouse. The fluids are pooled and frozen at -80.degree. C. After thawing the pool is centrifuged at 16.000 rpm for 30 minutes. The fat at the top is sucked off and the supernatant free of debris is saved. When necessary the centrifugation is repeated A crude immunoglobulin fraction is obtained from the ascitic fluid by 18% Na.sub.2 SO.sub.4 precipitation at room temperature. Subsequently this fraction is passed. through Sephacryl G 200 (Pharmacia) according to manufacturer's instructions using 0,1M Tris-HCI buffer pH 8.2. Active fractions are pooled and concentrated by Amicon XM 50 filters (Amicon). Protein determination is done by OD.sub.280 measurement assuming that 1 mg of provein gives rise to an absorbance of. . . .

ANSWER 10 OF 11 USPATFULL L19

Radiolabeling antibodies and other proteins with technetium or rhenium TΙ by regulated reduction

PΙ US 5078985 1992010/

This example illustrates the process of this invention for labeling DETD immunoglobulin G (IgG). 16. 3 obtained obtained from animals such as sheep, goats, mice or umans. Sodium Pertechnetate-Tc-99m U.S.P. is obtained. . . pot/ssium biphthalate and 10 mM sodium tartrate solution (pH 5.6) was a Jied '2 ml of 0.5 M stannous chloride in concentrated HCl (12 4). The Lannous chloride was prepared by adding the concentrated refer cloric acid to non-oxidized pellets of SnCl.st . . avin . surface free of dull stannous oxide. The A NaOH to adjust to the final pH. An IgG pH of the resultar preparation was me . : di' 0.25 al of Immune Globulin (Human), U.S.A., 3i. i.e. which contained 15-18% protein stabilized with 0.2000. The lycine, with 7.25 ml of Sterile Water for Injection, U.S.P., and ilt org through a 0.22 micron filter. 5 ml of the circ lution was mixed with 7.5 ml of the IgG preparation. It is ontaining the admixed. . . was collected and the remaining of rate, which contained the stannous and other salts, was distanded the reduced protein fraction was concentrated by training a concentration of 1.7 mg/ml. 0.5 ml. i.g. is of the concentration of 0.7 mg/ml. 0.5 ml. i.g. is of the concentration of 0.1 mg/ml. 0.5 ml. i.g. is of 0.1 mg/ml. serum i.g. and frozen. 0.5 ml of 0.1 mg/ml. SnCl.sub.2 prepared 1 40 mM potassium biphthalate/10 mM sodium tartrate solution, at pH 5.6, s added without allowing the reduced antibody solution to thaw, and is solution was also frozen.

A sterile, 3 mm diameter tin metal shot was added, the vial flushed with $N.\,\mathrm{sub.2}$ and stored at minus 20.degree.. . .

L19 ANSWER 11 OF 11 USPATFULL

Oligopeptides and intermediates and processes for their manufacture US 4720483 19880119

TI PI DETD

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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APPLICATION NO. DATE
    PATENT NO.
               KIND DATE
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    DE 19538625
                  A1
                       19970424
                                   DE 1995-19538625 19951017
    DE 19538625
                  C2 19990812
                 A1 19970424
                                   WO 1996-EP4476 19961015
   WO 9714717
       W: JP, US
       RW: AT, BE, CH, DE, DR, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
    EP 856008 A1 19980305
                                 EP 1996-934688 19961015
    EP 856008
                  В1
                       20010117
      R: DE, FR, GB
    JP_11514358 T2 19991207
                                   JP 1996-515514 19961015
   US 6001974
                  A 19991214
                                   US 1998-41082 19980310
PRIORITY APPLN: INFO.:
                                   DE 1995-19538625 19951017
                                   WO 1996-EP4476 19961015
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Albumin is sepd. and purified from human serum in a yield of AΒ .gtoreq.84% and a purity of .gtoreq.97% by successive chromatog. on a strongly basic anion-exchange membrane and a strongly acidic cation-exchange membrane. The albumin eluted in the 1st step can be applied to the membrane in the 2nd step without special conditioning. Thus, frozen serum was thawed at 4.degree., centrifuged at 15,000 g, filtered through a microfiltration membrane with pore size 0.45 .mu.m, desalted by gel filtration, adjusted to pH 5.4 to ppt. euglobulins, centrifuged at 15,000 g, frozen, thawed, centrifuged at 12,000 g, and adsorbed on anion-exchange membrane Sartobind Q 100; impurities were eluted with 25 mM NaOAc buffer (pH 5.4), and albumin was eluted with 50 mM NaOAc buffer (pH 4.5). The eluate was applied directly to cation-exchange membrane Sartobind S 100; impurities were eluted with 50 mM NaOAc buffer (pH 4.5), and albumin was eluted with 25 mM NaOAc buffer (pH 5.4) contg. 150 mM NaCl in 95% yield and 98% purity.

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L13 ANSWER 4 OF 11 HCAPLUS COPYRIGHT 2001 ACS >
ACCESSION NUMBER: 1985:209415 ECAPLUS
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DOCUMENT NUMBER: 102:20:419

Purified concentrate of the group TITLE:

G, A, M serum immunoglobulins

Step nek, Trun INVENTOR(S):

C ah. PATENT ASSIGNEE(S): SOURCE: 5 Ć. :ZX ···

DOCUMENT TYPE: ₽. . . . ze. LANGUAGE:

FAMILY ACC. NUM. TOU PATENT INFORMATION:

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PATENT NO. P. L. J. APPLICATION NO. DATE

CS 211239 B 1932c J CS 1980-4699 19800702

Aq. Iq solns, were cc 1. by s lionary freezing and thawing and used 15 to out get. Thus, at 2-5% soln, of conventional Iq pt. cd. rogen-free water was dialyzed against an aq. 13Cl 15th of 17 tred through get to remove low-mol swt. compds. This same as sterilized, gradually free.
                                                                                                                                                    _____
                                                                                                                                                    19800702
AB
            low-mol.-wt. compds. The s . was sterilized, gradually frozen in a metal column ar -56 dec. e. ... -5. degree. in 2-24 h, and
            thawed in 2-48 h at comp. thiq. conc. was batchwise collect. f. A . B comm. vit anal. co
                                                                                           The vit anal. control until the protein
                                                         at the same
            concn. dropped to
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Factor VIII with 25.4 times higher activity than that of the original plasma. Fibrinogen was pptd. by addn. of polyethylene glycol (PEG) and then sepd., and the Factor VIII was pptd. by addn. of more PEG. The pptd. Factor VIII was washed with 1.8-M glycocoll, dissolved in tris-Na citrate buffer, filter-sterilized, placed in dosage units, and freeze-dried. When the dried product was reconstituted to the same vol. with H2O, the soln. contained 25-30 Factor VIII units/mL (3 units/mg protein), and only 0.048 mg fibrinogen/mg protein.

L13 ANSWER 7 OF 11 HCAPLUS COPYRIGHT 2001 ACS ACCESSION NUMBER: 1973:513666 HCAPLUS

DOCUMENT NUMBER: 79:113666

TITLE: Bicchemical studies on cachexia due to cancer. I.

Partial purification and properties of the

hypoalbuminemic substance from Ehrlich solid carcinoma

AUTHOR(S): Kubota, Yukiho; Ueki, Hiroshi; Shoji, Shozo;

Shigematsu, Hidenari

CORPORATE SOURCE: Fac. Pharm. Sci., Kumamoto Univ., Kumamoto, Japan

SOURCE: Yakuyaku Zasshi (1973), 93(7), 887-92

CODEN: YKKZAJ

DOCUMENT TYPE: Journal

LANGUAGE: Japanese

A hypoalbuminemic substance was extd. from mouse Ehrlich solid carcinoma. The tumors were homogenized with saline, centrifuged, and the resultant turbid supernatant (F-1) was sepd. The turbidity was removed by freezing and thawing of the supernatant, and the resultant clear ext. (F-2) was dialyzed against 12mM NaCl. The inner soln. (possessing hypoalbuminemic activity) was freeze-dried. This fraction (F-3) was gel-filtered through Sephadex G-150, by using 12mM NaCl followed by M HOAc contg. 12mM NaCl as the eluant. An active fraction (F-4-II) was gel-filtered again under the same condition. Of 4 peaks obtained, the third was the active fraction (F-5-III). The yield in protein was 3.0% for F-1 and 0.2% for F-5-III. The decrease in serum albumin was statistically significant for F-1 and F-5-III, while the changes in serum globulins were not significant. F-5-III did not produce a significant change in total serum protein. F-5-III was not electrophoretically homogeneous, although it sedimented as a single peak in ultracentrifugal and . Its mol. wt. was roughly estd. to be 16,000 by ultracentrifugation. The amino acid content of F-5-III was 59.95%, and ... aspartic acid, glutamic acid, and isoleucine were present in fairly large amts. Neither the ext. prepd. from the gluteus maximus of a normal mouse by the same procedures nor bovine serum albumin had hypoalbuminemic activity.

L13 ANSWER 8 OF 11 HCAPLUS COPYRIGHT 2001 ACS

ACCESSION NUMBER: 1973:122436 HCAPLUS

DOCUMENT NUMBER: 73:122426

TITLE: Therapeutic concentrate of coagulation factors II, IX,

and X from citrated, factor VIII-depleted plasma Middleton, Sarah M.; Bennett, Ida H.; Smith, J. K. Scott. Natl. Blood Transfus. Assoc., R. Infirm.,

CORPORATE SOURCE: Scott. Natl. Blo Edinburgh, Scot.

SOURCE: Vox Sang. (1973), 24(5), 441-56

COLEN: VOSAAD

DOCUMENT TYPE: Journal LANGUAGE: English

AUTHOR(S):

AB DEAE-cellulose (1 kg, Whatman DE 32) was equilibrated with 0.5N HCl; water (to pH >4); 0.5N NaOH; water (to pH <8); 1.0N NaCl; water (to cond. <1 .mu.mho) and stirred for 9c min at 2.degree. with 120 l. of Cohn fraction I supernatant (from peoled ACD plasma) dild. with 40 l. of water. After centrifugation in a Sharples centrifuge, the supernatant was processed for IgG and albumin, while the absorbent was stirred with 41 (cooled to 5.degree.) of 0.03M LaH2PO4, 0.03M Na citrate, pH 6.9. cond. 9 mmho, ionic strength 0.24, and poured into a column of diam. 150-225 mm

(preautoclaved). The column was washed further with 4 l. of the same buffer and eluted with 6 l. of 0.2M NaCl in the same buffer (cond. 20 mM, ionic strength 0.44). Factors II, IX, and X, which emerged after the salt front, were collected in 400 ml fractions, assayed, pooled to 30 units of factor IX/ml, frozen, thawed, filter
-sterilized, and lyophilized in 10 ml doses. Reconstituted doses contained 0.6-1.2 g% protein, pH 6.8-7.1, 210-280m- M Ns, 40-90mM citrate, 20-50mM phosphate, 5-20mM Cl; cond. was 13 mmho, and osmolarity, 320 mosmolar. Factors II, IX, and X were 250-300-fold concd., recovered in 79% yield (range 42-105%). Although the material was pyrogenic in rabbits, this effect might have been due to heteroagglutinins; no pyr.genic reactions were obsd. in 30 patients receiving a total of 1200 doses at 30-50 units kg during hemorrhage, and 20-30 units/kg for mainte ance. The half-life of administered factor IX was 24-48 hr.

L13 ANSWER 9 OF 11 HCAPLUS COPYRIGHT 2001 ACS

ACCESSION NUMBER: 1973:14417 HCAPLUS

DOCUMENT NUMBER: 78:14417

TITLE: Antigen extracts of animal thymocytes and

specie-specific antithymocytic serums and

.gamma.-globulins

INVENTOR(S):
Goret, Pierre; Toma, Bernard; Salmon, Henri

PATENT ASSIGNEE(S): Laboratoires Albert Rolland

SOURCE: Ger. Offen., 28 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent LANGUAGE: German FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION: .

PATE	ENT NO.	KIN	D DATE	API	PLICATION NO.	DATE
DE 2	2205897	А	19720824	DE	1972-2205897	19720208
DE 2	2205897	C3	19800522			
DE 2	2205897	B2	19790830			
GB 1	1342391	A	19740103	GB	1971-4276	19710209
CH 5	538866	Α	19730831	CH	1972-1244	19720128
BE 7	778754	A1	1972053	BE	1972-113451	19720131
ES 3	399465	A.	197501	FS	1972-399465	19720203
CA 9	955849	Αï	19741011	CA	1972-134005	19720204
RO 6	61514	P	197612±5	RO	1972-69679	19720204
IL 3	38704	A1	1.3741 31	IL	1972-38704	19720207
DK 3	132104	В	19751017	DK	1972-527	19720207
NO 1	137308	В	19771 31	NO	1972-311	19720207
ZA 7	7200825	А	19721 .5	ZA	1972-825	19720208
FI 5	50203	В	3.750930	FΞ	1972-344	19720208
NL 7	7201683	Α	14/20511	. NL	1972-1683	19720209
AT 3	316743	В	15740 25	AT	1972-1039	19720209
PRIORITY	APPLN. I	NFO.:		GB	1971-4276	19710209

Thymus antigens with stabilar on long storage are injected into animals of species different than the antigen donor for the preph. of species-specific anti-thymocyles sera and gamma.-globulins. Cellular suspensions from the thymus or the bursa of fabricius of birds are filtered through a siness sieves or through gauze, frozen at -20 to -40.deg for the antigen and dispersed for use by thawing und a move and shough closely packed 51/8 mm diam. spherical code and the interest injected into animals of a species different than the antigen donor of a similar obtained from the liver and series of the antigen donor or a similar animal, and 5% by wt. glubarilds yde soln. gamma. Globulins are also prepd. from the tis surmocytic serum.

PATENT INFORMATION:

L13 ANSWER 1 OF 11 HCAPLUS COPYRIGHT 2001 ACS 1999:577986 HCAPLUS ACCESSION NUMBER: 132:34468 DOCUMENT NUMBER: Improved methods for the separation and TITLE: purification of immunoglobulin from egg yolk by filtration and one step anion-exchange chrc..atography Li, Jian-Cai; Chen, Tian-Bao; Zhang, Rong-Zhen; Chen, AUTHOR(S): Ru-Ming; Li, Long; Rao, Ping-Fan Institute of Biotechnology, Fuzhou University, Fujian, CORPORATE SOURCE: 350002, Peop. Rep. China Food Health Pac. Rim, Int. Conf. Food Sci. Technol., SOURCE: 3rd (1999), Meeting Date 1997, 384-390. Editor(s): Whitaker, John R. Food & Nutrition Press: Trumbull, Corin. CODEN: 68BQAF DOCUMENT TYPE: Conference LANGUAGE: English Reported in this paper is an improved method for isolation of antibodies by filtration and anion-exchange chromatog. from egg yolk. yolk was dild. 7 times with distd. water without pH adjustment and then frozen at -10.degree.C. The frozen dild. egg yolk was then thawed at 15.degree, C. Celite was added to the dild. egg yolk prior to filtration to facilitate the process. The dild. yolk soln. was then filtered through cellulose acetate filters with pore size of 0.22 um at 15.degree.C. An industrially feasible filtration velocity was achieved with the system, hundreds of times faster than reported data. A clear water-sol. fraction (WSF) contained approx. 15-18% by wt. of antibodies compared to the wt. of total proteins obtained. The WSF was then adjusted to contain 75 mM sodium phosphate buffer, pH 6.8, and applied to a DEAE-Toyopearl 650M column. electrophoretically pure IgY was obtained in the eluate eluted with 150 mM sodium phosphate buffer, pH 6.8. It is a method most likely to be developed into a com. scale process to produce IgY suitable for industrial applications in terms of cost and scale. REFERENCE COUNT: 19 (1) Akita, E; J Food Sci 1992, V57, P629 HCAPLUS REFERENCE(S): (2) Bucolo, G; Clin Chem 1973, V19, P476 HCAPLUS (3) Fichtali, J; Biotechnol Bioengin 1992, V40, P1388 HCAP1 US (5) Hatta, H; Agric Biol Chem 1990, V54, P2531 HCAPLUS (7) Horikoshi, T; J Food Sci 1993, V58(4), P739 HCAPLUS ALL CITATIONS AVAILABLE IN THE RE FORMAT L13 ANSWER 2 OF 11 HCAPLUS COPYRIGHT 2001 ACS 1399: /2617 HCAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 130:2:3644 Dried biologically or therapeutically active TITLE: preparations Kanellos, Jerry; Oates, Adrian; Goss, Neil INVENTOR(S): CSL Limited, Australia PATENT ASSIGNEE(S): PCT Int. Appl., 28 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT:

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PATENT NO. KIND DATE
                                              APPLICATION NO. DATE
     WO 9910011 A1 19990304 WO 1998-AU682 19980825
         W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG,
              KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
          UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES,
FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI,
              CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                                                  19980824
     ZA 9807633
                       A 19990225 ZA 1998-7633
                        A1 19990316 AU 1998-87231 19980825
A1 20000621 EP 1998-938550 19980825
     AU 9887231
     EP 1009438
            AT, BE, CH, DE, Dr. DS, rF GB, GR, IT, LI, LU, NL, SE, MC, PT,
              IE, FI
                                                                  19970825
                                                AU 1997-8719
PRIORITY APPLN. INFO.:
                                                WO 1998-AU682
                                                                  19980825
     A dried, heat-treated product comprises (i) a heat labile, biol. or
AB
     therapeutically active protein or peptide prepn. and (ii) a stabilizing
     effective amt. of a compn. comprising sucrose, trehalose and at least one
     amino acid. The protein or peptide prepn. may be, for example, a factor
     VIII conc. or a von Willebrand Factor conc. Fresh
     frozen plasma (FFP) is thawed at temps. below 5.degree.
     and the FVIII-rich cryoppt. is collected by centrifugation. The FVIII is
     extd. with Tris buffer. Levels of unwanted proteins, principally fibrinogen, fibronectin, Ig and albumin, are reduced by pptn.
     with heparin followed by repptn. of FVIII with sodium chloride/glycine
     buffer. The purified FVIII is redissolved in a sodium
     chloride-Tris-citrate bufter contg. sucrose and a low level of calcium
     chloride. The dissolved ppt. is filtered, treated with
     solvent/detergent and incubated. The mixt, is then filtered and
     chromatographed on a Sephacryl S400 column pre-equilibrated in the same
     buffer. The FVIII-rich eluate (>50 IU/mg total protein) is then
     concd. by ultrafiltration against the same buffer and chem.
     stabilizers added to the retentate. The bulk formulated conc.
     is sterile filtered, dispensed, freeze dried and heat
     treated at 80.degree. for /2 l. The freeze drying cycle
     proceeds under conditions of pagrammed temp./vacuum/timing for approx.
     100 h. The formulates product is loaded into a freeze dryer and
     the shelves cooled to -50.\deg r \cdot \epsilon.. The vacuum is applied and the temp.
     ramped up to -5 .degree.. The finished lyophilized product is then heated
     in a hot air oven at 80.degree. for 72 h.
REFERENCE COUNT:
                           10
                            (1, AJahi Crem Ind Co Ltd; JP 05331071 A 1993 HCAPLUS
REFERENCE(S):
                            (2) Asa ii > em Ind To Ltd; JP 06321805 A 1994 HCAPLUS
                           (3) religiorer. Aktiengesellschaft; US 4297344 A 1981
                                 CAPL .
                            (4) La ringwerk A tiengesellshaft; US 4562072 A 1985
                                 ..PLUS
                            . ) Et de la boratories Inc; US 4623717 A 1986 HCAPLUS
                           . L C .ATI: US AV. ILABLE IN THE RE FORMAT
L13 ANSWER 3 OF 11 HCAPLUS COLYRTHAT 2001 ACS
ACCESSION NUMBER:
                           199
                                 32145% HCAPLUS
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DOCUMENT NUMBER:
                            \ensuremath{\text{M}^{\prime\prime}} 'or for paration of albumin from serum by
TITLE:
                            i .e. ha c..comatography with membrane adsorbents
                           D shar: Wc ;ang; Nussbaumer, Dietmar; Kula,
Mc ia.egi ; Thoemmes, Joerg; Gebauer, Klaus-Heinrich
INVENTOR(S):
                            Sa orius Ag, Germany
PATENT ASSIGNEE(S):
                            Ge. Ofich., 5 pp.
SOURCE:
                            COD V: G XBX
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37

L13 ANSWER 10 OF 11 HCAPLUS COFFRIGHT 2001 ACS ACCESSION NUMBER: 1972::59953 HCAPLUS

DOCUMENT NUMBER: 77:59953

TITLE: A hypoalbuminemic substance from Ehrlich solid

carcinoma

AUTHOR(S): Kubota, Yukiho; Ueki, Hiroshi; Shigematsu, Hidenari CORPORATE SOURCE: Fac. Pharm. Sci., Kumamoto Univ., Kumamoto, Japan

SOURCE: Gamn (1972), 63(2), 277-8

COPEN: GANNA2

DOCUMENT TYPE: Journal LANGUAGE: English

The tumors were homogenized with saline and centrifuged at 15,000 rpm for 30 min; the turbid supernatant (F-1) was sepd., the turbidity was removed by repeated freezing and thawing followed by centrifugation, and the resulting clear supernatant (F-2) was dialyzed against 12 mM NaCl soln. The inner soln. possessing hypoalbuminemic activity was freeze-dried. This fraction (F-3) was gel-filtered through Sephadex G-150, using 12 mM NaCl soln, as eluant. Of the 4 peaks obtained, the 3rd was the active fraction (F-4III). For its assay, the samples were given to dd-K mice by a daily i.p. injection for 6 days. Serum albumin and qlobulins were detd. by densitometry of the electrophoregram of the serum on cellulose acetate paper. A decrease in serum albumin was statistically significant for F-1 and F-4-III, while the changes in total serum protein or serum globulins were not significant. The hydrolysate of F-4-III consisted of hexose, hexosamine, and 18 kinds of amino acids, in which aspartic acid, glutamic acid, and isoleucine were contained in fairly large amts. (>10% each). Neither the ext. prepd. from gluteus maximus muscle of a normal mouse by the same procedures nor bovine serum albumin displayed hypoalbuminemic activity.

L13 ANSWER 11 OF 11 HCAPLUS COFYRIGHT 2001 ACS

ACCESSION NUMBER: 1967:8300 HCAPLUS

DOCUMENT NUMBER: 66:9300

TITLE: Preparation of purified antihemophilic globulin and a study of its properties

AUTHOR(S): Rozenberg, G. Ya.; Rutberg, R. A.; Novoselova, V. N.;

Blinova, M. A.; Papush, N. D.

CORPORATE SOURCE: Central list. Hematol. and Blood Transfusion, Moscow,

USSR

SOURCE: Probl. Gematol. Pereliv. Krovi (1966), 11(10), 3-8

CODEN: PGPKA8

DOCUMENT TYPE: Journal LANGUAGE: Russian

Mix fresh human blood (450 ml.) with 50 ml. 3.8% di-Na citrate and 0.003% levomycetin (1:9), centrifuge twice at 4.degree. and 1600-1800 rpm. for 20 and 30 min., resp. Freeze the plasma immediately at -35.degree. and keep at this temp. for 1-2 weeks. Thaw it at + 10.degree. for 1-3.3 hrs., decrease its temp. (2-3.degree.) to 0.degree., and centrifuge at this temp. for 40 min. and 1800 rpm. Use the supernatant, which is free from the main part of antihemophilic globulin (AHG), for the prepn. of fibrinogen (I) and other protein prepns. Dissolve the ppt. from 1 1.. of plasma in 80 ml. of distd. apyrogenic water (total vol. about 90 ml.), add 10 ml. sterile Al(OH)3 gel, mix vigorously, and let stand for 15-18 i.rs. Centrifuge at 1800 rpm. for 40 min. at O.degree., sep. the supernatant, add 20 ml., i.e., 2% of the initial vol. of buffer soln. (21.25 g. NaCl, 12.5 g. glucose, and 5 g. di-Na citrate in 500 ml. water, pH 6.5-6.7), and filter through a plastic filter, freeze, and lyophilize. One 1. of blood plasma yielded 100-110 ml. AHG soln. To purify it, take the supernatant after Al(OH)3 removal, add slowly and dropwise EtOH to final concn. of 20% (the temp. must not exceed 0.degree.), let stand 1 hr., and centrifuge at 0.degree. and 1800 rpm. for 40 min. Dissolve the ppt. in buffer solm. (0.2 g. di-Na citrate, 0.8 g. NaCl, and 0.5 g.

glucose in 100 ml. water, pH 6.5–6.7) (10% vol. of the initial plasma), and lyophilize. A 12–15–fold enrichment of AHG was reached. The crude prepn. contained about 250 mg. % I, albumin <250 mg. %, but pptn. with EtOH removed it (simultaneously a 30% decrease of AHG activity occurred). Since no prepn. was pyrogenic or toxic (tested on rabbits, 1 ml./kg., and mice, 0.5 ml./mouse), 50–100 ml. of the prepn. was given 5 times to 4 patients with hemophilia, with successful results.

=> d his 17-(FILE 'HCAPLUS' ENTERED AT 10:25:54 ON 08 MAR 2001) E E58+OLD E IMMUNO GLOBULIN/CT E IMMUNOGLOBULIN/CT E E93+ALL E E103+ALL E IMMUNOGLOBULINS/CT E E107+ALI 190743 S ?GLOBULIN? OR POLYGLOBIN? OR IG OR IG## OR E138-E154 OR BENCE L7 72406 S FREEZE? OR FROZ? L8 L9 14925 S THAW? 238456 S FILTER? L10 2001179 S CONCEN? L117 S L7 AND L8 AND L9 AND L10 AND L11 L12 11 S L7(L)(ISOL? OR PURIF? OR L11) AND L8 AND L9 AND L10 L13 => file uspatfull COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 127.86 171.86 TOTAL DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE ENTRY SESSION -12.35-12.94 CA SUBSCRIBER PRICE FILE 'USPATFULL' ENTERED : 10:52:5. ON 08 MAR 2001 CA INDEXING COPYRIGHT (C) 2 J3 AMEL CAN CHEMICAL SOCIETY (ACS) FILE COVERS 1971 TO PATTENT FUBLICATION DATE: 6 Mar 2001 (20010306/PD) FILE LAST UPDATED: 6 Mar 1001 (21310306/ED) HIGHEST PATENT NUMBER: US6199207. CA INDEXING IS CURRENT THROUGH 6 M : 2001 (20010306/UPCA) ISSUE CLASS FIELDS (/INCL) CUFRENT THOOUGH: 6 Mar 2001 (20010306/PD) REVISED CLASS FIELDS (/NCL) LAST RE O. JED: Dec 2000 USPTO MANUAL OF CLASSIFICATIONS THE AURUS ISSUE DATE: Oct 2000 >>> Page images are availab. For at hts from 1/1/1997. Current >>> week patent text is typically liked by Thursday morning and >>> page images are available for dislay by the end of the day. >>> Image data for the /FA Firly .re vai able the following week. <<< <<< <<< <<< >>> Complete CA file indexing on closical ratents (or equivalents) <<< >>> is included in file product Products is available for the <<< >>> USPTO Manual of Constitution of the constitut

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PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH
FIELD CODE - 'AND' OPERATOR ASSUMED 'L7(P) (ISOL?'
PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH
FIELD CODE - 'AND' OPERATOR ASSUMED 'L11) (P) L8'
PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH
FIELD CODE - 'AND' OPERATOR ASSUMED 'L8(P)L9'
PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH
FIELD CODE - 'AND' OPERATOR ASSUMED 'L9(P)L10'
         21952 ?GLOBULIN?
             0 POLYGLOBIN?
          9218 IG
           332 IGS
          9406 IG
                  (IG OR IG.
         32336 IG##
            70 IGA/CT
             5 IGA1/CT
             3 IGA2/CT
             8 IGD/CT
            62 IGE/CT
           233 IGG/CT
             O "IGG AUTOANTIBODIES"/CT
             7 "IGG CONJUGATES"/CT
            82 IGG1/CT
            16 IGG2/CT
            20 IGG2A/CT
             9 IGG2B/CT
            12 IGG3/CT
            11 IGG4/CT
            94 IGM/CT
             O "IGM AUTOANTIBODIES"/CT
             3 IGY/CT
           134 BENCE
             1 BENCES
           135 BENCE
                  (BENCE OR BENCES)
         18603 JONES
        109513 PROTEIN -
           108 BENCE-JONES PYOTEIN?
                  (BENCE (W) JONES (W) PROTEIN?)
        390807 ISOL?
        210332 PURIF?
        748745 CONCEN?
         60200 FREEZE?
         51396 FROZ?
         20426 THAW?
        546264 FTTF
          3344 L*(P (IS * : ), PURTO OR (L1) (F) 8(P) L9(P) L10
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PROXIMITY OPERATOR LEVEL NOT CONTEN WITH
FIELD CODE - 'AND' CPER OF AS 12 3) L9'
PROXIMITY OPERATOR LEVEL HOL CC 2 TEL WITH
FIELD CODE - 'AND' OPER.TOR SS. WI > 'L9(S)L10'
         21952 ?GLOBULI: ?
             O POLYGLOBIL '
          9218 IG
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10 11 Nov. 10 15

ACCESSION NUMBER:

1982:488019 HCAPLUS

DOCUMENT NUMBER:

97:88019

TITLE:

Isolation and purification of

human thyroglobulin

AUTHOR(S):

Lu, Fengxian; Yang, Cuipo; Tong, Zhigang; Tang, Te;

Znang, Tiangeng; Wang, Renming; Li, Yan

CORPORATE SOURCE:

Dep. Pathol., Tianjin Med. Coll., Tianjin, Peop. Rep.

China

SOURCE:

Tianji. Yiyao (1980), 8(1), 8-11

CODEN: TIYADG

DOCUMENT TYPE:

Journa.

LANGUAGE:

Ch. Les

Human thyroid tissue was . zen at -20.degree., thawed , sectioned, and 100 g of : cti ned tissue was treated with 300 mL saline contg. a few drops of $0.3 \, e^{-p_h M \varepsilon}$ and kept in a refrigerator overnight. The ext. was filtered, concd., ...d _entrifuged at 4000 rpm for 30 min to give a supe latent which was passed through a Sephadex G 200 column. A fraction contg. thyroglobulin was collected, concd., and again passed through a Sephadex G 200 column with a saline eluent. The active fraction was collected, distributed into ampuls, and freeze dried. A yield of 1.5 g thyroglobulin/100 g tissue was obtained. The product was purified further by ion-exchange chromatog. The sedimentation coeff. (S20,w) was 14.0 for thyroglobulin from deceased normal subjects but was 14.3 for that from patients with hyperthyroidism.

L13 ANSWER 6 OF 11 HCAPLUS COPYRIGHT 2001 ACS

ACCESSION NUMBER:

1978:27788 HCAPLUS

DOCUMENT NUMBER:

88:27788

TITLE:

Purified antihemophilic globulin A

(factor-VIII)

INVENTOR(S):

Iga, Yoshiro; Shiga, Masashi G. en Cr ss Corp., Japan

PATENT ASSIGNEE(S):

Gc . Offen., 21 pp.

SOURCE: COL N. GWXXBX

DOCUMENT TYPE:

Pater.

LANGUAGE:

AΒ

Cerman

FAMILY ACC. NUM. COURT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2715832	A1	1977 (1)	DE 1977-2715832	19770407
DE 2715832	B2	19793"ld		
DE 2715832	C3	19 00.00	•	
JP 52125609	A2	19 71 11	CP 1976-39894	19760409
JP 55012890	34	10900 04		
US 4093608	F.	1′78 -(US 1977-783625	19770401
PRIORITY APPLN. INFO.			JP 1976-39894	19760409

Purified antihem, hile: glob lin A (1 stor VIII [9001-27-8]) is obtained from fresh or pooled human plasma by use of a weakly basic (diethyla inc'ethvi dextran (DEAE dextran) ion exchanger. The stability and soly of the actor VIII are also increased and the prothrombin complex and f.: inc. in ontents are decreased using this method. For example, i end from a man plasma (contg. 1.3 .times. 106 units to at , ot. ombin activity and 1.3 .times. 106 units total Factor VIII activity. . . thamed, pooled, adjusted to 4.degree., pH 7.4, and the preswollen (pH 7.1 site of the plant of the plant of the proting of the plant of the plant was entrifuged to sep. Fact VI as a cryc pr. The ppt. was extd. with a tris-Na citrate buff 1 (ph 1.4) give 1 crude soln. contg. a 75.3% yield of

1.7.1

and the same of

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16 IGG2/CT
    20 IGG2A/CT
    9 IGG2B/CT
    12 IGG3/CT
    11 IGG4/CT
    94 IGM/CT
     O "IGM AUTOANTIBODIES"/CT
     3 IGY/CT
   134 BENCE
     1 BENCES
   135 BENCE
         (BENCE OR BENCES)
18603 JONES
109513 PROTEIN?
   108 BENCE-JONES PROTEIN?
         (BENCE (W) JONES (W) PROTEIN?)
 60200 FREEZE?
 51396 FROZ?
 20426 THAW?
546264 FILTER?
748745 CONCEN?
  3294 L7(S)L8(S)L9(S)L10(S)L11
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=> d hit

L16

L16 ANSWER 1 OF 3294 USPATFULL

One of the preferred methods for introducing DNA into a plant cell by SUMM means of Agrobacterium is the so-called leaf disk transformation using Agrobacterium [Horsch et al (1985)]. Sterile leaf disks from a suitable, target plant are incubated with Agrobacterium cells comprising one of the GA 20-oxidase-encoding DNA sequence according to the invention, and are then transferred into or onto a suitable nutrient medium. Especially suitable, and therefore preferred within the scope of this invention, are LS media that have been solidified by the addition of agar and enriched with one or more of the plant growth regulators customarily used, especially those selected from the group of the auxins consisting. of a-naphthylacetic acid, picloram, 2,4,5-trichlorophenoxyacetic acid, 2,4-dichlorophenoxyacetic acid, indole-3-butyric acid, indole-3-lactic acid, indole-3-succinic acid, indole-3-acetic acid and p-chlorophenoxyacetic acid, and from the group of the cytokinins consisting of kinetin. 6benzyladenine, 2-isopentenyladenine and zeatin. The preferred concentration of auxins and cytokinins is in the range of from 0.1 mg/l to 10 mg/l.

DETD 50 to 500 mg/ml of DNA fragments are added to a reaction batch in the buffer recommended by the manufacturer, New England Biolabs. The reaction batch contains all four deoxynucleotide triphosphates in concentrations of 0.2 mM. The reaction takes place over a period of 30 minutes at 15 C and is then terminated by heating at 65 C for 10 minutes. For fragments obtained by cleaving with restriction endonucleases that produce 5'-projecting ends, such as EcoRI and BamHI, the large fragment, or Klenow fragment, of DNA polymerase is used. For fragments obtained by means of endonucleases that produce 3'-projecting ends, such as PstI and Sacd, the T4 DNA polymerase is used. The use of these two enzymes is described on pages 113 to 121 of the Maniatis et al (1982) reference.

The extracted DNA is first treated with restriction enzymes, then subjected to electro-phoresis in a 0.8% to 1% agarose gel, transferred to a nitrocellulose membrane (Southern E. M. (1975)] and hybridised with the DNA to be detected which has previously been subjected to nick-translation (DNA-specific activities of 5.times.10.sup.8 to 10.times.10.sup.8 c.p.m/mg). The filters are washed three times for 1 hour each time with an aqueous solution of 0.03 M sodium citrate and 0.3 M sodium chloride at 65 C. The hybridised DNA is made

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332 IGS
            9406 IG
                     (IG OR IGS)
           32336 IG##
              70 IGA/CT
               5 IGA1/CT
               3 IGA2/CT
               8 IGD/CT
              62 IGE/CT
             233 IGG/CT
               O "IGG AUTOANTIBODIES"/CT
               7 "IGG CONJUGATES"/CT
              82 IGG1/CT
              16 IGG2/CT
              20 IGG2A/CT
               9 IGG2B/CT
              12 IGG3/CT
              11 IGG4/CT
              94 IGM/CT
               O "IGM AUTOANTIBODIES"/Ci
               3 IGY/CT
             134 BENCE
               1 BENCES
             135 BENCE
                     (BENCE OR BENCES)
           18603 JONES
         109513 PROTEIN?
             108 BENCE-JONES PROTEIN?
                     (BENCE (W) JONES (W) PROTEIN?)
         390807 ISOL?
         210332 PURIF?
         748745 CONCEN?
           60200 FREEZE?
           51396 FROZ?
           20426 THAW:
         546264 FILTER?
            3344 L7(S'(ISOL? OR PULIF? OR L11)(S)L8(S)L9(S)L10
L15
=> s 17(s)(isol? or pu. if: 21 + 111)(:,1.(s)19(s)110
\Rightarrow s 17(s)18(s)19(s 11 (s' 11
PROXIMITY OPERATOR LEVIL NO CONLISTENT WITH FIELD CODE - 'AND' OPERATO ASSUL D 'L7(S) L8'
PROXIMITY OPERATOR LEVEL N CONS GENT WITH FIELD CODE - 'AND' OPERATOR SUM' 'L8(S) 59' PROXIMITY OPERATOR LEVEL NG! CONS "ENT WITH
FIELD CODE - 'AND' OPERATOR ASSU' : 9(S)L10'
PROXIMITY OPERATOR LEVEL NOT CON 311 (F WITH
FIELD CODE - 'AND' OPERATOR ASSU & D 10(S)L11'
           21952 ?GLOBULIN?
               O POLYGLOBIE
            9218 IG
             332 IGS
            9406 IG
                     (IG OR TGS
           32336 IG##
              70 IGA/CT
                5 IGA1/CT
                3 IGA2/CT
               8 IGD/CT
               62 IGE/CT
             233 IGG/CT
                O "IGG AUTOANT1BODIES"/CT
                7 "IGG CONJUGATES"/CT.
              82 IGG1/CT
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\Rightarrow s ?globulin?(s)18(s)19(s)110(s)111
          21952 ?GLOBULIN?
          60200 FREEZE?
          51396 FROZ?
          20426 THAW?
         546264 FILTER?
         748745 CONCEN?
              11 ?GLOBULIN? (S) L8 (S) L9 (S) L10 (S) L11
L19
=> d ti pn kwic tot
L19 ANSWER 1 OF 11 USPATFULL
        Process for the manufacture of thrombin inhibitors
TΙ
        US 5728549 19980317
PΙ
          . . one week, 2 to 5.times.10.sup.6 cloned hybridoma cells are
DETD
        injected intraperitone lly. Ascillo fluid is repeatedly taken from each
        mouse and frozen at -80 degase. C. The accumulated fluid is
        thawed and centrifuged for 30 minutes at 4.degree. C. and 16,000
        revs/min. The fat is filtered off with suction and 0.9 volume
        equivalents of a saturated ammonium sulphate solution is slowly added
        dropwise, while stirring at 0.degree. C., to the remaining debris-free
        supernatant. The resulting crude immunoglobulin fraction is
        passed through Sephacryl G 2000 (Pharmacia) using 0.1 mM tris.HCl (pH
        8.2), in accordance with the instructions of the manufacturer. Active
        fractions are combined and concentrated with Amicon XM50
        filter (Amicon).
L19 ANSWER 2 OF 11 USPATFULL
        Cryptosporidium hybrid vector and transformed host cells
TТ
        US 5643772 19970701
PΙ
        For SDS-PAGE, 2.times.10.sup.9 oocysts were lysed by 5 cycles of
DETD
        freeze-thawing in 1% Triton Buffer (150 mM Na CI; 100
        mM EDTA; and 1% Triton X-100), in the presence of protease.
        hrs. (Petersen, C., at al., "Characterization of an Mr>900,000
        Cryptosporidium parvum Sporozoites Glycoprotein Recognized by
        Hyperimmune Bovino Cc'ostral Immunoglobulin", Inf. & Immun.
        60(12):5132 (1932)). Eastern blots were incubated with HBC Ig (lot
        \#40529) (dil 1/500) \therefore 20 ml PBS. . . "Cryptosporidium parvum
        (Apicomplexa: Cryptosporidiidae) Oocyst and Sporozoite Antigens
        Recognized by Bovine Colostral Antibodies", Inf. Imm. 58:2966 (1990)).
        Eluted antibodies were filter sterilized and
        concentrated to a final volume of 1 ml in a Centriprep 10
        concentrator (Amicon, Mass.)
L19 ANSWER 3 OF 11 JSPATFOLL
        Direct radiolabeling of antik lies and other proteins with technetium or
ΤI
        rhenium
PΙ
        US 35500 19970506
        US 5102990 19920407 ( rigin...)
       An IgG preparation was the billion of Immune Globulin (Human), I'm the billion of Immune 15-18% protein staking the billion of Ith 1.21-0.32M glycine, with 7.25 ml of Sterile Water for the U.T., and filtering through a 0.22 micros filter. I the I (II) reducing solution was mixed with 7.5 ml of the I (II) reducing solution was mixed with 7.5 ml of the I (II) and other salts, was discarded. The reduced and Sn (II) completed protein fraction was concentrated by ultrafiltration to a concentration of 1.7 mg/ml 0.5 mg
DETD
        by ultrafiltration to a concentration of 1.7 mg/ml. 0.5 mg
        aliquots of reduced and small) complexed protein were placed in sealed,
        N.sub.2 gas filled . . . . vi and frozen. A Sn (II) pertechnetate reduci . sol. on was made of 0.5 ml of 0.1 mM SnCl.sub.2
        in 40 mM potassium b. hthatate/ J. . . at a pH of 5.6. The Sn (II)
        pertechnetate reducin schitio, was added without allowing the reduced
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antibody solution to thaw, and this solution was also frozen. A sterile, 3 mm diameter tin metal shot was added, the vial flushed with N.sub.2 and stored at -20.degree. C....

- L19 ANSWER 4 OF 11 USPATFULL . The
- TI Radiolabeling antibodies and other proteins with technetium or rhenium by regulated reduction
- PI US 35457 19970218 : US 5078985 19920107 (Original) : 1
- This example illustrates the process of this invention for labeling DETD immunoglobulin G (IgG). IgG is obtained from animals such a sheep, goats, mice or humans. Sodium Pertechnetate-Tc-99m U.S.P. is obtained from. . . potæssium biphthalate and 10 mM sodium tartrate solution (pH 5.6) was add.d 0.2 ml of 0.5 M stannous chloride in concentrated HCl (12 M). The stannous chloride was prepared by adding the concentrated hydrochloric acid to non-oxidized pellets of SnCl.sub.2 having a surface free of dull stannous oxide. The pH of the resultant. .. 1 M NaOH to adjust to the final pH. An IgG preparation was made by diluting 0.25 ml of Immune Globulin (Human), U.S.P., Cutter Biological, which contained 15-18% protein stabilized with 0.21-0.32 M glycine, with 7.25 ml of Sterile Water for Injection. U.S.P., and filtering through a 0.22 micron filter. 5 ml of the reducing solution was mixed with 7.5 ml of the IgG preparation. The vial containing the admixed. . . was collected and the remaining eluate, which contained the stannous and other salts, was discarded. The reduced protein fraction was concentrated by ultrafiltration to a concentration of 1.7 mg/ml. 0.5 mg aliquots of reduced protein were placed in sealed, N.sub.2 gas filled serum vials and frozen. 0.5 ml of 0.1 mM SnCl.sub.2 prepared in 40 mM potassium biphthalate/10 mM sodium tartraté solution, at pH 5.6 was added without allowing the reduced antibody solution to thaw, and this solution was also frozen. A sterile, 3 mm diameter tin metal shot was added, the vial flushed with N.sub.2 and stored at m us 20.degree.. .
- L19 ANSWER 5 OF 11 SPECIFIE ?
- TI Method of producing an anci- immunoglobulin concentrate and a pharmaceutical preparation
- With the method according to the invention, immunoglobulin can be obtained directly from human plasma. The starting material is the plasma from Rh-neg. The denors sensitized to the rhesus factor D. The individual don tion cotains through plasmapheresis is preferably frozen and carrifully thave at O.Cagree. A. degree. C. before fracticiation, and oled. The plasma pool should contain more than 10 .mu.g anti-D Ir perml. The cation exchange chromatography calber rised reliminarily using any known method, such as, for example, and so refer editing the cryoglobulins or by fractionated precipitation by respectively and the cryoglobulins or by fractionated precipitation by the results of ethanol (Cohn, E. J. et al. J. Am. Chem. Soc. 68, 45t. And the plasma precipitation of the immunoglobulis of animals of the cryoglobulin substance of the immunoglobulis of animals of such methods, such as precipitation of the immunoglobulis of animals of such methods, such as precipitation of the immunoglobulis of animals of such methods. Prior to the cation exchange chromatography according to the invention, the cryoglobulin effect plasma or the presman section containing immunoglobulin dissolved in the equilibration is buffer is preferably filtered, and to inactivate his den vires at it is treated with a biocompatible, organic solvent such as the containing immunoglobulin determined to the cation that the containing immunoglobulin determined to the cation that the containing immunoglobulin determined to the cation containing immunogl

92.

being, as a. . . detergents used for virus inactivation are likewise removed here. The anti-D IgG of the IgG subclasses 1 and 3 specifically concentrated by this process step is eluted by the ion exchange gel by increasing conductivity to preferably over 10 mS/cm. The. less than 15% of the total IgG fraction and has a changed IgG subclass spectrum: IgG1 and IgG3 are strongly concentrated, IgG2 and IgG4 are greatly reduced. This anti-D IgG fraction is then purified further by treatment with a second, weakly. . . diethylaminoethyl (DEAE) as functional $grou_P$, anti-D IgG not being bound to the ion exchangers under the selected conditions. To increase concentration, the purified anti-D IgG fraction is preferably bound a second time to a CM ion exchange gel and is eluted as concentrated as possible under suitable conditions, and is manufactured into the "ine" of Juct. For achieving the conditions according to the invertion of a not of significance whether the anti-D IgG fraction is elute; by no. asing the ionic concentration, or by pH shift or by a lover that yet in composition. To reduce the concentration of further underired components, such as, for example, proteases, the immunoglobulin solution can be treated additionally with an adsorbent, such as, for example, aluminum hydroxide gel, during any stage of the. . .

- ANSWER 6 OF 11 USPATFULL L19
- Process for the manufacture of thrombin inhibitors TI
- US 5422249 1995060€ PΙ
- . . . one week, 2 to 5.times.10.sup.6 cloned hybridoma cells are DETD injected intraperitoneally. Ascitic fluid is repeatedly taken from each w. mouse and frozen at -80.degree. C. The accumulated fluid is thawed and centrifuged for 30 minutes at 4.degree. C. and 16,000 revs/min. The fat is filtered off with suction and 0.9 volume equivalents of a saturated ammonium sulphate solution is slowly added dropwise, while stirring at 0.degree. C. to the remaining debris-free supernatant. The resulting crude immunoglobulin fraction is passed through Saphacryl 2000 (Pharmacia) using 0.1 mM tris.HCl (pH 8.2), in accordance with the instructions of the manufacturer. Active fractions are combined and concentrated with Amicon XM50 filter (Amicon).
- L19 ANSWER 7 OF 11 USPATFUL:
- Direct radiolabeling of substrates containing monosulfides or disulfide bonds with radionuclides TΤ
- US 5277893 19 10111 PΙ
- An IgG prepara on as ..ad by diluting 0.25 ml of Immune DETD Globulin (Human), 6 3.3 , eatter Biological, which contained 15-18% protein Flabilized with 0.21-0.32 M glycine, with 7.25 mi of Sterile Water for Injectic. U.S.P., and filtering through a 0.22 micron filter. 5 . of the Sn (II) reducing solution was mixed with 7.5 ml of the Ich preparation. The vial containing. . . contained the Sn (II), and other salts, was discarded. The reduced and Sn (I) and the root in fraction was concentrated by ultrafiltation of reduced and Sn (I) complexed protein were placed in sealed, N.sub.2 gas fill ... vials and frozen. A Sn (II) pertechnetate reducing oluti was made of 0.5 ml of 0.1 Mm SnCl.sub.2 in 40 mM potassium biphthale 10. . . at a pH of 5.6. The Sn (11) pertechnetate reducing . 10.10.1 was added without allowing the reduced antibody solution to them, and this solution was also frozen. A sterile. I import tin metal shot was added, the vial flushed with ... id + ored at -20.degree. C.. . .
- L19 ANSWER 8 OF 11 USPAL LL
- Direct radiolabelia of ...til dies and other proteins with technetium or TIrhenium

n v av.

visible by blackening an X-ray film over a period of 24 to 48 hours. An oligo(dT)-primed cDNA library was constructed in .lambda.gtll DETD (Amersham) using cotyledon mRNA. The total library (70,000 clones) was amplified to give 3.5.times.10.sup.8 plaque forming units (pfu)/ml of which 69% were recombinant. Immunoscreening of the amplified library was performed with 3,600 pfu on one 90 mm plate and probing with the 20-oxidase peptide autibody (1 mg/ml) and an alkaline phosphatase-conjugated anti-rabbit IgG second antibody. When the protein was incubated at increasing concentrations DETD with [.sup.14 C] GA.sub.1? (Table 1A) sequential oxidation of the C-20 methyl group to the alcohol, aldehyde and carboxylic acid occurred to give, respectively, radiolabelled GA.sub.15, GA.sub.24 and GA.sub.25 as products. The corresponding 13-hydroxy GA products (GA.sub.44, GA.sub.19 and GA.sub.17) were also obtained, although at lower efficiency, when the lysate was incubated wit' [.sup.14 C] GA.sub.53 (Table 1B). A comparison of the aldehyge substrates GA.sub.24 (non-hydroxylated), GA.sub.19 (monohydro.tyle ad) and GA.sub.23 (dihydroxylated), showed that the efficiency of oxidation to the corresponding tricarboxylic acids decreased with increasing polarity of the substrate (Table 1C). In addition, the corresponding C.sub.19 "GA products (GA.sub.9, GA.sub.20 and GA.sub.1), which are formed by loss of C-20 as CO.sub.2, were obtained in low yield. The results indicate that a single enzyme may catalyse each of the steps involving oxidation at C-20 during GA biosynthesis, possibly also including the loss of C-20, although confirmation of this must awaic studies with the corresponding enzyme from a plant tissue in which C.sub.19 -GA production forms a major Explants roughly 5 to 10 mm are cut from young leaves 3 to 5 cm long and DETD third to sixth from the apex of N. tabacum cv 'Xanthi nc' grown under axenic conditions [Facciotti and Pilet, 1979] in solid MS medium [Murashige and Skoog, 1962] containing 0.7% phytagar (Gibco-BRL), 1 mg/1IAA, 0.15~mg/l kinetin. These explants are plated on solid MS medium containing 0.6% phytagar, 40 mg/l adenine sulfate, 2 mg/l IAA, and 2 mg/l kinetin on the sur ace of which is placed a #1 Whatman filter and incubated for 24 hr in the dark at 24 C. Agrobacterium strains containing the binary vectors described above are grown overnight in LBMG at 30 C on a shaker at 180 rpm. Explants are dipped into a pacturial suppression of 3.3.times.10.sup.8 cells/ml for approximately 5 minutes, 110 led on sterile paper towels, and re-plated. on the same plates. After 48 hours explants are placed on selection medium containing the same paste medium as above plus 350 mg/l cefotaxime and 100 ag/l lanar/cip. Co-cultivated control tissue is placed on the same redire but without kanamycin. The explants are transferred to fresh med a ev :y two weeks. Shoots are harvested 4 to 8 weeks after co-cultivation placed on 50 ml culture tubes with 25 ml of solid MS medium co. This 0.% hytogar, 1 mg/l IBA, 350 mg/l cefotaxime, and 10t mg/l kanarycin. All tissue is grown at 24 C to 28 C, 12 hours of light, 12 hours dark, light intensity 6700 to 8400 1 .times.. Shoots root it 1 to 2 weeks and are then transplanted to planting mix in 4" para and paced in the "transgenic plant phytotron". 1 to 1.5 ml PCV of the suspendion culture cells from above are incubated DETD in 10 to 15 ml of a filter sterilized mixture consisting of 4% cellulase RS with 1% Rhc syst in KMC (8.65 g/l KCl, 16.47 g/l MgCl.sub.2.6H ..b 0 a. 112... g.1 CaCl.sub.2.2H.sub.2 0, pH 5.6) salt solution. Dig tion is at ied it at 30 C on a slow rocking table for a period of 3 to 4 hours. We preparation is monitored under an inverted microscope for problem in the protoplasts which are released are collected as for ...: The preparation is filture hrough a 100 mm mesh sieve, DETD followed by a 50 mm $_{\odot}$:si $_{\odot}$. .he protoplasts are washed through the sieves with a volume of $_{\odot}4C$. It solution equal to the original volume sieves with a volume of MC lt solution equal to the original volum of enzyme solution. 10 ml of the protoplast preparation is placed in each of several disposable pristic centrifuge tubes, and 1.5 to 2 ml of 0.6 M sucrose solution (buffered to pH 5.6 with 0.1% MES and KOH)

layered underneath. The tube is centrifuged at 60 to 100.times.g for 10 minutes, and the protoplasts banding at the interface collected using a pipette and placed in a fresh tube. The protoplast preparation is resuspended in 10 ml of fresh KMC salt solution, and centrifuged for five minutes at 60 to 100.times.g. The supernatant is removed and discarded, and the protoplasts resuspended gently in the drop remaining, and then 10 ml of a 13/14 strength KMC solution gradually added. After centrifuging again for five minutes, the supernatant is again removed and the protoplasts resuspended in a 6/7 strength KMC solution. An aliquot is taken for counting, and the protoplasts again sedimented by centrifugation. The protoplasts are resuspended at 10.sup.7 per ml in KM-8p medium or in 0.5 M mannitol containing 6 mM MgCl.sub.2 or other suitable medium for use in transformation as described in the following examples. This protoplast suspension is used for transformation and is cultured as described helo

DETD A. The protoplasts of recognished at the last step of above in a 0.5 M mannitol solution or main. The 12 to 30 mM MgCl.sub.2. A heat shock of 45.degree. C. for file must as is given as describes The protoplasts are distributed in aliquets to transfer mation in centrifuge tubes, 0.3 ml of suspended protoplas a per tube. During the next 10 minutes the following are added: D.A and PEG solution (MW 6000, 40% containing 0.1 M Ca(NO.sub.3).sub.2 and 0.4 M mannitol; pH 8 to 9 with KOH) to give a final concentration of 20% PEG. The aliquots are incubated for 30 minutes with occasional gentle shaking, and then the protoplasts are placed in petri dishes (0 ml original protoplast suspension per 6 cm diameter dish) and cultur is a described.

DETD C. The above is repeated with the modification that the final concentration of PEG is bet meen 13 and 25%.

DETD

DETD

Protoplasts are prepared from embryogenic suspension cultures of above by aseptically filtering the cells on a Nalgene 0.2 mm filter unit and then adding 0.5 g fresh weight cells to each 12.5 ml of protoplasis enz me mixture in a petri dish. The enzyme mixture consists of the elithase RS, 7 mM CaCl.sub.2 xH.sub.2 O, 0.7 mM NaH.sub.2 PO4xH.sub. C, 3 mM MES (pH 5.6), glucose (550 mOs/kg H.sub.2 O of pH 5.6), and is filter terilized. The mixture is swirled on an orbital sharer a above 50 r m in dim (<420 l.times.) light for about 4 to 5 hours. The dige is then sieved through a stainless steel sieve (100 mm mesh size) and istributed into 12 ml centrifuge tubes which are centrifuged at abou. 60 to 100.times.g for about 5 minutes. The protoplast-containing rediment is then washed three times with protoplast culture medium KM-79 adjusted to 550 mOs/kg H.sub.2 O with glucose. At this point a flotation step may be included for further purification of the proton standard lathis case, the washed protoplasts are layered atop 10 ml. or of -8p culture medium adjusted to 700 mOs/kg H.sub.2 O wit sucrese. .. contribugation at 60 to 100.times.g for about 10 minu is, to top . I be diag at the interface are collected using a fine pipette. First the protoplasts are resuspended in 1 to 2 ml KM-8p culture medit as eved through a stainless steel screen (20 medium suitable for afofotic according to the examples below.

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A. The purified recoplate te, ated at a density of about 5.times.10.sup. Lotoplate terml in KM-8p culture medium containing 1.3% SeaPlaque a glose (El C.p., Marine Colloids Division, Rockland, Me., USA) and 3 to 40° for aditioned medium (obtained from 3 to 4 week-old Dactylis glossata a embryogenic suspension cultures by filtering the medium scraph a sterile Nalgene 0.2 mm filter, making the modum standard sterile Nalgene 0.2 mm filter, making the modum standard sterile Nalgene 0.2 mm filter, making the modum standard sterile Nalgene 0.2 mm filter, making the modum standard sterile Nalgene 0.2 mm filter, making the modum standard sterile Nalgene 0.2 mm filter, making the modum standard sterile Nalgene 0.2 mm filter, making the modum standard sterile Nalgene 0.2 mm filter ster

50 rpm in light at 670 1.times.. New suspension cultures are formed as the colonies grow out of the agarose and release cells into the liquid medium. The resultant suspension cultured cells are plated onto agar-solidified SH-30 medium and placed in the dark at 25.degree. C. until callus is formed.

A. Immediately after purification of the protoplasts, electroporation is DETD performed according to Shilito et al (1985) using linezed plasmid The protoplasts are resuspended after the last wash at a density of about 7.times.10.sup.6 protoplasts per ml in the electroporation buffer (0.4 M $\,$ mannitol, 6 mM MgCl.sub 2). The protoplasts are placed in 0.7 ml aliquots in 10 ml plastic centrifuge tubes. Plasmid DNA and sonicated calf thymus DNA (Sigma) to give final concentrations of 10 mg/ml and 50 mg/ml respectively is added to the tubes. Then 0.38 ml PEG solution [24% PEG 6000 in 0.4 M mannitol, 30 mM MgCl.sub.2, 0.1% MES (H 5.6)] is added and the sclution gently mixed. The protoplast suspension is transferred into the charber of a Dialog Electroporator and 10 pulses of 3250 Vcm.sup.-1 initial voltage and exponential decay constant of 10 msec applied at 30 sec inter als. The sample is removed from the chamber, and placed in a 10 cm diameter petri dish. 10 ml of KM-8p medium containing 1.2% SedFlaque agarose is added, the protoplasts distributed evenly throughout the medium, and the agarose allowed to gel.

The cells are given a plasmolysis treatment before bombardment. Packed cell volume is measured and colls are diluted in 1 MS liquid medium with added osmoticum: 0.4 M sorbitol for suspension cells and 0.6 M sorbitol for callus cells. Cells are diluted such that the final packed cell volume per target is 1/20 ml for a fine suspension and 1/10 ml for callus. Diluted cells are placed in a 250 ml flask containing a stir bar and are stirred for a minimum of 30 minutes, up to a few hours. To plate the cells, 2 ml is withdrawn from the flask and pipetted into the top of a vacuum flask onto which a Whatman 2.5 cm GFA filter has been placed. The vacuum is applied until the cells are dried onto the filter, the filters are placed on 60.times.15 mm petri plates containing 5 ml of solid post bombardment plasmolysis medium: lMS containing 0.2 M sorbitol for suspension cells, or 0.4 M sorbitol for callus cells. Two filters are plated on each dish.

Bombarding of cell cultures is carried out using a device as described in EP-A... Invest the petri plate containing the cell filters onto the platform on top of the stage, centered over the particle flight opening. In the clear lid over the top of the platform. Place a microproject le onto the breech pin and close the breech. Push the "arm" by ton to fill the reservoir with the appropriate amount of helium gas insully 1800-1900 psi). Pull the vacuum on the chamber to sup, about 12 mm. Turn off the vacuum, and push the "arm" and "fire buth 13. Mo at the "arm" buttor on the "off" position. Each filter is usually she two.

filter is usually shelf two sections and the dark overnight. The next day, filters are readily if modes a post-bombardment for suspension cells a readily if 7-10 days post-bombardment for suspension cells a readily for callus cells. Cells are scraped off the filters are given to the surface of plates containing 1MS plus and a property of the selection marker generated in plant transformation. Plates are incubated in the dark for sevent weeks. Resistant colonies that arise after a few weeks are transferre of 1M. reselection agent. Coloni that continue to proliferate for about 3-4 deks are then transferred to "0.5 MS" maintenance medium: 4M. so the date of the selection agent. Coloni that continue to proliferate for about 3-4 deks are then transferred to "0.5 MS" maintenance medium: 4M. so the date onto this medium biweekly until embryogenic structures of it we seems suitable for regeneration.

DETD Seeds of Arabidopsis that and are being erecta are surface sterilised by treatment with 5% the chlorite solution in 0.01% Tween-20 (Sigma), washed to exite the call and suspended in 0.15% agar. The seeds

treatment with 5% : Schlorite solution in 0.01% Tween-20 (Sigma), washed to exit, er and suspended in 0.15% agar. The seeds are sown onto 0.8% par core in ing Murashige and Skoog Medium supplemented with sigma; (Sigma) and 5% sucrose in sterile Magenta

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by autoradiography against Kodak X-OMAT AR film with intensifying screens, overnight at -70.degree. C. The insert of pAt2204 's abelled with .sup.32 P-dCTP and used to probe DETD nitrocellulose filter lifts of a full-length cDNA library, constructed in .lamaia.ZapII (Stratagene) from poly-A.sup.4 RNA isolated from shoot material of the gibbereffin-deficient gal mutant of Arabidopsis thaliana (Koornneef M and van der Veen J H (1980)). The hybridisation is carried out in 50% fromamide, 50 mM sodium phosphate pH 6.3, 0.75 M NaCl, 75 mM sodium citrate, 0.1% bovine serum albumin, 0.1% Ficoll 400, 0.1% polyvinylpyrrolidone 360, 0.1% sodium dodecyl sulphate and 100 .mu.g/ml salmon testes DNA at 42.degree. C. overnight Filters are washed in 15 mM NaCl, 1.5 mM sodium citrate at 42.degree. C. for 10 min. Hybridising plaques are identified by autoradiography and purified by successive rounds of hybridisation. Positive clones are converted into pBluescript clones by plasmid rescue and characterised by FraRI digestion and DNA sequencing. Clone pAt2353 and clone pAt2301, containing a 1.3 kbp insert, are chosen for heterologous expression

The 1.3 kbp insert p/
gel electrophoresis and ated to expression vector pTrcHisA DETD (Invitrogen), previously cap with EquRI and dephosphorylated. Ligation products are introduced into E. coli strain TOP10 (Invitrogen) by transformation and selected by growth on 2.times. YT agar with ampicillin at 100 .mu.g/ml. Plasmid DNA is isolated from a number of the resulting clones and the orientation of the cDNA insert determined by HindIII digestion. Clone pAt2328 contained a cDNA insert in sense orientation, and is used to inoculate 50 ml of 2.times. YT containing ... carbenicillin at 100 .mu.g/ml. After 2 hrs growth shaking at 37.degree. C., IPIG (Isopropyl-b-D-thiogalactopyranoside) is added to 1 mM and the cultures are grown for a further 5 hrs. The cells are collected by centrifugation and suspended in 4 ml of 100 mM Tris-Cl pH7.5, 4 mM DTT and sonicated on ice for a total of 90 secs. The samples are then frozen in liquid nitrogen, thawed by hand and insoluble material removed by centrifugation at 15,000.times.q for 5 min. The resulting surequatant material is stored at -80.degree. C. and t subsequently used for nayme assay. <-----User Break--- --> => s 17(p)18(p)19(p)1.0(p).11PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH FIELD CODE - 'AND' OPERATO' AS JMI'D 'L7(P) L8' PROXIMITY OPERATOR LEVEL NOT C NSI TENT WITH FIELD CODE - 'AND' OPL' & Cr AS .MF '1 (P)19' PROXIMITY OPERATOR LEV ... CO... ' TE WITH FIELD CODE - 'AND' OPERATOR ASSU 1 1/(P)L10' PROXIMITY OPERATOR LEVEL NOT CO STATE OF WITH FIELD CODE - 'AND' OPERATOR ASSEMENT 'LIC(P)L11' 21952 ?GLOBULIN? O POLYGLOBIN? 9218 IG 332 IGS 9406 IG , (G O), 10 32336 IG## · 70 IGA/CT 5 IGA1/(..

3 IGA2/CT 8 IGD/CT

62 IGE/CT

233 IGG/CT

O "IGG AUTO" TIBODIF "/CF

containers (Sigma). Plants are grown for 4 weeks at 20.degree. C. and shoot material is frozen in liquid nitrogen and stored at -70 degree. C. Genomic DNA is isolated essentially as described by Murray and Thompson (Murray MG and Thompson WF (1980)). The frozen tissue, 10 g, is ground to a slurry in an ice-cooled mortar with a small amoust of acid-washed sand. The homogenate is transferred to a polypropylene centrifuge tube and an equal volume of 2% (w/v) CTAB (cetyltimethy ammonium bromide, Sigma), 1.4M NaCl, 0.1 M Tris-Cl pH8.0, 20 mM EDIA added . After gentle mixing, the tube is incubated at 67.degree. C. for 20 min with occasional mixing. The tube is removed from the water bath and 0.5 volumes of chloroform added, mixed gently and left at room temperature (20.degree. C.) for 20 min with occasional inversion. The tube is centrifuged at 2000 g for 5 min at room temperature, the upper phase removed to a new tube and the lower phase discarded. To the uppe phase is added 0.1 vol of 10% (w/v) CTAB, 0.7M NaCp and the chiorogram extraction above is repeated. The upper phase is again decanted () a new tube and 2 volumes of 1% (w/v) CTAB, 50 mM Tris-Cl, pH8.0, 10 mM EDTA added. This is mixed gently and left at room temperature for 1 hour, then centrifuged at 5000 mg for 5 min. The pellet is dissolved in 50% (w/w) CsCu in TB, buffer with ethidium bromide at 0.5 mg/mid. The solution is transferred to a Quick-seal tube (Beckman) and centrifuge in a vertical rotor (Beckman VTi90) for 16 hours at 80,000 rpm at 20 degree. C. The DNA is visualised under natural l ight and removed with t e aid of a syringe and nee. Ethidium bromide is removed by extraction four times with 5 volumes of butan-1-ol, previously equilibrated against NaCI-saturated water. The solution is diluted by the addition of 3 volumes of TB buffer (10 mM_{i} Tris-Cl pH8.0, 1 mM EDTA) and DNA precipitated with 2 volumes of EtOH. The DNA is pelleted by centrifugation at 10,000 g for 10 min at O.degree. C., washed with 70% ETOH, dried in vacuo, and dissolved in TE buffer. The DNA concentration is determined by its absorbance at 260 nm.

DETD A 50 ml aliquot of 2.times. YT (1.6% Bactotryptone, 1% yeast extract, 0.5% NaCl) including 0.2% maltose and 10 mM MgSO.sub.4 is inoculated with a single colony of E coli XL1-Blue. This is gown overnight at 30 C, transferred to a sterile centrifuge tube and spun down at 2000.times.g for 5 min, room temperature. The cells are resuspended in 10 Mm MgSO.sub.4. In steril 15 ml tubes, 500 ml E. coli cells is mixed with 50,000 recombinant interphage from the amplified library and incubated at at room temperature for 10 min followed by 37 C for 15 min. Molten top agarest 10 75 in 2.times. YT/0.2% maltose/10 mM MgSO.sub.4), 6.5 ml, is add in the continuity of a plate of 1.5% agar in 2.times. YT/0.2% maltose/10 mM MgSO.sub.4. The plates are incubated inverted at 37 C for 6 hours and then store involve. ight at 4.degree. C. Duplicate nitrocellulose filters in the are air dried and treated for 5 min each in 1.5 n. Jan 0.5 m NaOH (denaturation); 3M NaCl, 1M Tris-Cl pH 6.5 (new fail in the continuity of the continuity of the paper and baked in vacuo, lower new are wetted in Nature of filter paper, at 80 C for 2 hrs.

DETED Nitrocallulose filters if the propered as above are wetted in Nature of the contents of the paper.

Nitrocellulose filter ift prepared as above are wetted in water and prehybrid e for 2 hours at 42.degree. C. in hybridization buffer (50% formain: 50 NaPi pH 6.3, 0.75 M NaCl, 75 mM trisodium citrate, 0.1% (w/v) in turn bumin, 0.1% (w/v) Ficoll 400, 0.1% (w/v) polyvinylpyrrox ion 1% //v) sodium dodecyl sulphate (SDS), 100 mg/ml sonic if sill. The probe is boiled for 2 min, mixed with 25 m ori ic fer and sealed into a polythene bag with the piehyb in if it in it and sealed into a polythene bag with the piehyb in if it is discontant probe is removed by washing in 0.3 M NaCl, 30 mM tris discontant citrate, 0.1% (w/v) SDS at room temperature for 15 min, and in 1 Mil 21, 1.5 mM trisodium citrate, 0.1% SDS at 60.degree. C. for 2 mlr. I sitively-hybridizing plaques are identified

The second secon

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7 "IGG CONJUGATES"/CT
            82 IGG1/CT
            16 IGG2/CT
            20 IGG2A/CT
            9 IGG2B/CT
            12 IGG3/CT
            11 IGG4/CT
            94 IGM/CT
             0 "IGM AUTOANTIBODIES"/CT
             3 IGY/CT
           134 BENCE
             1 BENCES
           135 BENCE
                 (BENCE OR BENCE:)
         18603 JONES
        109513 PROTEIN?
           108 BENCE-JONES PROTE N?
                 (BENCE(W) JONE?(W) Pt ""EIN?)
         60200 FREEZE?
         51396 FROZ?
         20426 THAW?
        546264 FILTER?
        748745 CONCEN?
L17
          3294 L7(P)L8(P)L9(P)L10(P)L11
\Rightarrow s ?golulin?(s)18(s)19(s)110(s)111
             0 ?GOLULIN?
         60200 FREEZE?
         51396 FROZ?
         20426 THAW?
        546264 FILTER?
        748745 CONCEN?
L18
             0 ?GOLULIN?(S)' *(') ±9(S) L10(S) L11
=> s ?globulin?(s)18(s)19(s)1.0(s)111
         21952 ?GLOBULIN?
         60200 FREEZE?
         51396 FROZ?
         20426 THAW?
        546264 FILTER,
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CS Dep. Pharm. Therap., Univ. Sheffield, Sheffield, Engl.

SO Clin. Sci. (1966), 31(2), 215-21

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